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(54) **Phagemid-based method of producing filamentous bacteriophage particles displaying antibody molecules and the corresponding bacteriophage particles**

Methode zur Herstellung von filamentösen, antikörper-präsentierenden Bakteriophagenpartikeln durch Verwendung von Phagemiden und die dadurch hergestellten Bakteriophagenpartikel

Utilisation de phagemids dans une méthode de production de particules de bacteriophages filamenteux présentant des anticorps à leurs surfaces et les particules de bacteriophages correspondantes

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AL: "GENERATION OF A LARGE
COMBINATORIAL LIBRARY OF THE
IMMUNOGLOBULIN REPERTOIRE IN PHAGE
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Description

[0001] The present invention relates to methods for producing members of specific binding pairs. The present invention also relates to the biological binding molecules produced by these methods.

[0002] Owing to their high specificity for a given antigen, the advent of monoclonal antibodies (Kohler, G. and Milstein C; 1975 Nature 256: 495) represented a significant technical break-through with important consequences both scientifically and commercially.

[0003] Monoclonal antibodies are traditionally made by establishing an immortal mammalian cell line which is derived from a single immunoglobulin producing cell secreting one form of a biologically functional antibody molecule with a particular specificity. Because the antibody-secreting mammalian cell line is immortal, the characteristics of the antibody are reproducible from batch to batch. The key properties of monoclonal antibodies are their specificity for a particular antigen and the reproducibility with which they can be manufactured.

[0004] Structurally, the simplest antibody (IgG) comprises four polypeptide chains, two heavy (H) chains and two light (L) chains inter-connected by disulphide bonds (see figure 1). The light chains exist in two distinct forms called kappa (K) and lambda (λ). Each chain has a constant region (C) and a variable region (V). Each chain is organized into a series of domains. The light chains have two domains, corresponding to the C region and the other to the V region. The heavy chains have four domains, one corresponding to the V region and three domains (1, 2 and 3) in the C region. The antibody has two arms (each arm being a Fab region), each of which has a VL and a VH region associated with each other. It is this pair of V regions (VL and VH) that differ from one antibody to another (owing to amino acid sequence variations), and which together are responsible for recognising the antigen and providing an antigen binding site (ABS). In even more detail, each V region is made up from three complementarity determining regions (CDR) separated by four framework regions (FR). The CDR's are the most variable part of the variable regions, and they perform the critical antigen binding function. The CDR regions are derived from many potential germ line sequences via a complex process involving recombination, mutation and selection.

[0005] It has been shown that the function of binding antigens can be performed by fragments of a whole antibody. Example binding fragments are (i) the Fab fragment consisting of the VL, VH, CL and CH1 domains; (ii) the Fd fragment consisting of the VH and CH1 domains; (iii) the Fv fragment consisting of the VL and VH domains of a single arm of an antibody, (iv) the dAb fragment (Ward, E.S. et al., Nature 341, 544-546 (1989) which consists of a VH domain; (v) isolated CDR regions; and (vi) F(ab')₂ fragments, a bivalent fragment comprising two Fab fragments linked by a disulphide bridge at the hinge region.

[0006] Although the two domains of the Fv fragment are coded for by separate genes, it has proved possible to make a synthetic linker that enables them to be made as a single protein chain (known as single chain Fv (scFv); Bird, R.E. et al., Science 242, 423-426 (1988) Huston, J.S. et al., Proc. Natl. Acad. Sci., USA 85, 5879-5883 (1988)) by recombinant methods. These scFv fragments were assembled from genes from monoclonals that had been previously isolated. In this application, the applicants describe a process to assemble scFv fragments from VH and VL domains that are not part of an antibody that has been previously isolated.

[0007] Whilst monoclonal antibodies, their fragments and derivatives have been enormously advantageous, there are nevertheless a number of limitations associated with them.

[0008] Firstly, the therapeutic applications of monoclonal antibodies produced by human immortal cell lines holds great promise for the treatment of a wide range of diseases (Clinical Applications of Monoclonal Antibodies. Edited by S. S. Lennox. British Medical Bulletin 1984. Publishers Churchill Livingstone). Unfortunately, immortal antibody-producing human cell lines are very difficult to establish and they give low yields of antibody (approximately 1 μ g/ml). In contrast, equivalent rodent cell lines yield high amounts of antibody (approximately 100 μ g/ml). However, the repeated administration of these foreign rodent proteins to humans can lead to harmful hypersensitivity reactions. In the main therefore, these rodent-derived monoclonal antibodies have limited therapeutic use.

[0009] Secondly, a key aspect in the isolation of monoclonal antibodies is how many different clones of antibody producing cells with different specificities, can be practically established and sampled compared to how many theoretically need to be sampled in order to isolate a cell producing antibody with the desired specificity characteristics (Milstein, C., Royal Soc. Croonian Lecture, Proc. R. Soc. London B. 239; 1-16, (1990)). For example, the number of different specificities expressed at any one time by lymphocytes of the murine immune system is thought to be approximately 10⁷ and this is only a small proportion of the potential repertoire of specificities. However, during the isolation of a typical antibody producing cell with a desired specificity, the investigator is only able to sample 10³ to 10⁴ individual specificities. The problem is worse in the human, where one has approximately 10¹² lymphocyte specificities, with the limitation on sampling of 10³ or 10⁴ remaining.

[0010] This problem has been alleviated to some extent in laboratory animals by the use of immunisation regimes. Thus, where one wants to produce monoclonal antibodies having a specificity against a particular epitope, an animal is immunised with an immunogen expressing that epitope. The animal will then mount an immune response against the immunogen and there will be a proliferation of lymphocytes which have specificity against the epitope. Owing to

this proliferation of lymphocytes with the desired specificity, it becomes easier to detect them in the sampling procedure. However, this approach is not successful in all cases, as a suitable immunogen may not be available. Furthermore, where one wants to produce human monoclonal antibodies (eg for therapeutic administration as previously discussed), such an approach is not practically, or ethically, feasible.

[0011] In the last few years, these problems have in part, been addressed by the application of recombinant DNA methods to the isolation and production of e.g. antibodies and fragments of antibodies with antigen binding ability, in bacteria such as *E.coli*.

[0012] This simple substitution of immortalised cells with bacterial cells as the 'factory', considerably simplifies procedures for preparing large amounts of binding molecules. Furthermore, a recombinant production system allows scope for producing tailor-made antibodies and fragments thereof. For example, it is possible to produce chimaeric molecules with new combinations of binding and effector functions, humanised antibodies (e.g. murine variable regions combined with human constant domains or murine-antibody CDRs grafted onto a human FR) and novel antigen-binding molecules. Furthermore, the use of polymerase chain reaction (PCR) amplification (Saiki, R.K., et al., *Science* **239**, 487-491 (1988)) to isolate antibody producing sequences from cells (e.g. hybridomas and B cells) has great potential for speeding up the timescale under which specificities can be isolated. Amplified VH and VL genes are cloned directly into vectors for expression in bacteria or mammalian cells (Orlandi, R., et al., 1989, *Proc. Natl. Acad. Sci., USA* **86**, 3833-3837; Ward, E.S., et al., 1989 *supra*; Larrick, J.W., et al., 1989, *Biochem. Biophys. Res. Commun.* **160**, 1250-1255; Sastry, L. et al., 1989, *Proc. Natl. Acad. Sci., USA*, **86**, 5728-5732). Soluble antibody fragments secreted from bacteria are then screened for binding activities.

[0013] However, like the production system based upon immortalised cells, the recombinant production system still suffers from the selection problems previously discussed and therefore relies on animal immunization to increase the proportion of cells with desired specificity. Furthermore, some of these techniques can exacerbate the screening problems. For example, large separate H and L chain libraries have been produced from immunized mice and combined together in a random combinatorial manner prior to screening (Huse, W.D. et al., 1989, *Science* **246**, 1275-1281, WO90/14443; WO90/14424 and WO90/14430). Crucially however, the information held within each cell, namely the original pairing of one L chain with one H chain, is lost. This loses some, of the advantage gained by using immunization protocols in the animal. Currently, only libraries derived from single VH domains (dAbs; Ward, E.S., et al., 1989. *supra*.) do not suffer this drawback. However, because not all antibody VH domains are capable of binding antigen, more have to be screened. In addition, the problem of directly screening many different specificities in prokaryotes remains to be solved.

[0014] Thus, there is a need for a screening system which ameliorates or overcomes one or more of the above or other problems. The ideal system would allow the sampling of very large numbers of specificities (eg 10^6 and higher), rapid sorting at each cloning round, and rapid transfer of the genetic material coding for the binding molecule from one stage of the production process, to the next stage.

[0015] The most attractive candidates for this type of screening, would be prokaryotic organisms (because they grow quickly, are relatively simple to manipulate and because large numbers of clones can be created) which express and display at their surface a functional binding domain eg. an antibody, receptor, enzyme etc. In the UK patent GB 2137631B methods for the co-expression in a single host cell of the variable H and L chain genes of immunoglobulins were disclosed. However, the protein was expressed intracellularly and was insoluble. Further, the protein required extensive processing to generate antibody fragments with binding activity and this generated material with only a fraction of the binding activity expected for antibody fragments at this concentration. It has already been shown that antibody fragments can be secreted through bacterial membranes with the appropriate signal peptide (Skerra, A. and Pluckthun, A. 1988 *Science* **240** 1038-1040; Better, M et al 1988, *Science* **240** 1041-1043) with a consequent increase in the binding activity of antibody fragments. These methods require screening of individual clones for binding activity in the same way as do mouse monoclonal antibodies.

[0016] It has not been shown however, how a functional binding domain eg an antibody, antibody fragment, receptor, enzyme etc can be held on the bacterial surface in a configuration which allows sampling of say its antigen binding properties and selection for clones with desirable properties. In large part, this is because the bacterial surface is a complex structure, and in the gram-negative organisms there is an outer wall which further complicates the position. Further, it has not been shown that eg an antibody domain will fold correctly when expressed as a fusion with a surface protein of bacteria or bacteriophage.

[0017] Bacteriophage are attractive prokaryote related organisms for this type of screening. In general, their surface is a relatively simple structure, they can be grown easily in large numbers, they are amenable to the practical handling involved in many potential mass screening programmes, and they carry genetic information for their own synthesis within a small, simple package. The difficulty has been to practically solve the problem of how to use bacteriophages in this manner. A Genex Corporation patent application number WO88/06630 has proposed that the bacteriophage lambda would be a suitable vehicle for the expression of antibody molecules, but they do not provide a teaching which enables the general idea to be carried out. For example WO88/06630 does not demonstrate that any sequences: (a)

have been expressed as a fusion with gene V; (b) have been expressed on the surface of lambda; and (c) have been expressed so that the protein retains biological activity. Furthermore there is no teaching on how to screen for suitable fusions. Also, since the lambda virions are assembled within the cell, the fusion protein would be expressed intracellularly and would be predicted to be inactive. Bass et al., in December 1990 (after the earliest priority date for the present application) describe deleting part of gene III of the filamentous bacteriophage M13 and inserting the coding sequence for human growth hormone (hGH) into the N-terminal site of the gene. The growth hormone displayed by M13 was shown to be functional. (Bass, S., et al. *Proteins, Structure, Function and Genetics* (1990) 8: 309-314). A functional copy of gene III was always present in addition, when this fusion was expressed. A Protein Engineering Corporation patent application WO90/02809 proposes the insertion of the coding sequence for bovine pancreatic trypsin inhibitor (BPTI) into gene VIII of M13. However, the proposal was not shown to be operative. For example, there is no demonstration of the expression of BPTI sequences as fusions with protein VIII and display on the surface of M13. Furthermore this document teaches that when a fusion is made with gene III, it is necessary to use a second synthetic copy of gene III, so that some unaltered gene III protein will be present. The embodiments of the present application do not do this. In embodiments where phagemid is rescued with M13K07 gene III deletion phage, there is no unaltered gene III present.

[0018] WO90/02809 also teaches that phagemids that do not contain the full genome of M13 and require rescue by coinfection with helper phage are not suitable for these purposes because coinfection could lead to recombination.

[0019] In all embodiments where the present applicants have used phagemids, they have used a helper phage and the only sequences derived from filamentous bacteriophage in the phagemids are the origin of replication and gene III sequences.

[0020] WO90/02809 also teaches that their process needed information such as nucleotide sequence of the starting molecule and its three-dimensioned structure. The use of a pre-existing repertoire of binding molecules to select for a binding member, such as is disclosed herein, for example using an immunoglobulin gene repertoire of animals, was not disclosed. Further, they do not discuss favouring variegation of their binding molecules in natural blocks of variation such as CDRs of immunoglobulins, in order to favour generation of improved molecules and prevent unfavourable variations. WO90/02809 also specifically excluded the application of their process to the production of scFv molecules.

[0021] In each of the above discussed patents (WO88/06630 and WO90/02809), the protein proposed for display is a single polypeptide chain. There is no disclosure of a method for the display of a dimeric molecule by expression of one monomer as a fusion with a capsid protein and the other protein in a free form.

[0022] WO91/17271, citable under Article 54(3) EPC and only as entitled to its earlier claimed priority date, describes insertion of a protein into a coat protein of a bacteriophage and use of affinity purification techniques to select phage particles.

[0023] WO92/06204, also citable under Article 54(3) EPC only as entitled to its claimed priority date, proposes display of heteromeric receptors proteins on the surface of cells, citing filamentous bacteriophage as a preferred embodiment.

[0024] Another disclosure published in May 1991 (after the earliest priority date for the present application) describes the insertion into gene VIII of M13, the coding sequences for one of the two chains of the Fab portion of an antibody with co-expression of the other from a plasmid. The two chains were demonstrated as being expressed as a functional Fab fragment on the surface of the phage (Kang A.S. et al., (1991) *Proc. Natl. Acad. Sci. USA*, 88 p4363-4366). No disclosure was made of the site of insertion into gene VIII and the assay for pAb binding activity by ELISA used a reagent specific for antibody L chain rather than for phage. A further disclosure published in March 1991 (after the earliest priority date for the present application) describes the insertion of a fragment of the AIDS virus protein gag into the N-terminal portion of gene III of the bacteriophage fd. The expression of the gag protein fragment was detected by immunological methods, but it was not shown whether or not the protein was expressed in a functional form (Tsunetsugu-Yokota Y et al. (1991) *Gene* 99 p261-265).

[0025] The problem of how to use bacteriophages in this way is in fact a difficult one. The protein must be inserted into the phage in such a way that the integrity of the phage coat is not undermined, and the protein itself should be functional retaining its biological activity with respect to antigen binding. Thus, where the protein of choice is an antibody, it should fold efficiently and correctly and be presented for antigen binding. Solving the problem for antibody molecules and fragments would also provide a general method for any biomolecule which is a member of a specific binding pair e.g. receptor molecules and enzymes.

[0026] Surprisingly, the applicants have been able to construct a bacteriophage that expresses and displays at its surface a large biologically functional binding molecule (eg antibody fragments, and enzymes and receptors) and which remains intact and infectious. The applicants have called the structure which comprises a virus particle and a binding molecule displayed at the viral surface a 'package'. Where the binding molecule is an antibody, an antibody derivative or fragment, or a domain that is homologous to an immunoglobulin domain, the applicants call the package a 'phage antibody' (pAb). However, except where the context demands otherwise, where the term phage antibody is used generally, it should also be interpreted as referring to any package comprising a virus particle and a biologically functional binding molecule displayed at the viral surface.

[0027] pAbs have a range of applications in selecting antibody genes encoding antigen binding activities. For example, pAbs could be used for the cloning and rescue of hybridomas (Orlandi, R., et al (1989) PNAS 86 p3833-3837), and in the screening of large combinatorial libraries (such as found in Huse, W.D. et al., 1989, Science 246, 1275-1281). In particular, rounds of selection using pAbs may help in rescuing the higher affinity antibodies from the latter libraries. It may be preferable to screen small libraries derived from antigen-selected cells (Casali, P., et al., (1986) Science 234 p476-479) to rescue the original VH/VL pairs comprising the Fv region of an antibody. The use of pAbs may also allow the construction of entirely synthetic antibodies. Furthermore, antibodies may be made which have some synthetic sequences e.g. CDRs, and some naturally derived sequences. For example, V-gene repertoires could be made in vitro by combining un-rearranged V genes, with D and J segments. Libraries of pAbs could then be selected by binding to antigen, hypermutated in vitro in the antigen-binding loops or V domain framework regions, and subjected to further rounds of selection and mutagenesis.

[0028] As previously discussed, separate H and L chain libraries lose the original pairing between the chains. It is difficult to make and screen a large enough library for a particularly advantageous combination of H and L chains.

[0029] For example, in a mouse there are approximately 10^7 possible H chains and 10^7 possible L chains. Therefore, there are 10^{14} possible combinations of H and L chains, and to test for anything like this number of combinations one would have to create and screen a library of about 10^{14} clones. This has not previously been a practical possibility.

[0030] The present invention provides a number of approaches which ameliorate this problem.

[0031] In a first approach, (a random combinatorial approach, see examples 16 and 17) as large a library as is practically possible is created which expresses as many of the 10^{14} potential combinations as possible. However, by virtue of the expression of the H and L chains on the surface of the phage, it is reasonably practicable to select the desired combination, from all the generated combinations by affinity techniques (see later for description of selection formats).

[0032] In a second approach (called a dual combinatorial approach by the present applicants, see example 22), a large library is created from two smaller libraries for selection of the desired combination. This ameliorates the problems still further. The approach involves the creation of: (i) a first library of say 10^7 e.g. H chains which are displayed on a bacteriophage (as a fusion with the protein encoded by gene III) which is resistant to e.g. tetracycline; and (ii) a second library of say 10^7 e.g. L chains in which the coding sequences for these light chains are within a plasmid vector containing an origin of replication for a bacteriophage (a phagemid) which is resistant to e.g. ampicillin (i.e. a different antibiotic) and are expressed in the periplasmic space of a host bacterium. The first library is then used to infect the bacteria containing the second library to provide 10^{14} combinations of H and L chains on the surface of the resulting phage in the bacterial supernatant.

[0033] The advantage of this approach is that two separate libraries of eg 10^7 are created in order to produce 10^{14} combinations. Creating a 10^7 library is a practical possibility.

[0034] The 10^{14} combinations are then subjected to selection (see later for description of selection formats) as disclosed by the present application. This selection will then produce a population of phages displaying a particular combination of H and L chains having the desired specificity. The phages selected however, will only contain DNA encoding one partner of the paired H and L chains (deriving from either the phage or phagemid). The sample eluate containing the population is then divided into two portions. A first portion is grown on e.g. tetracycline plates to select those bacteriophage containing DNA encoding H chains which are involved in the desired antigen binding. A second portion is grown on e.g. ampicillin plates to select those bacteriophage containing phagemid DNA encoding L chains which are involved in the desired antigen binding. A set of colonies from individually isolated clones e.g. from the tetracycline plates are then used to infect specific colonies e.g. from the ampicillin plates. This results in bacteriophage expressing specific combinations of H and L chains which can then be assayed for antigen binding.

[0035] In a third approach (called a hierarchical dual combinational approach by the present applicants), an individual colony from either the H or L chain clone selected by growth on the antibiotic plates, is used to infect a complete library of clones encoding the other chain (H or L). Selection is as described above. This favours isolation of the most favourable combination.

[0036] In a fourth approach (called a hierarchical approach by the present applicants, see examples 18 and 36) both chains are cloned into the same vector. However, one of the chains which is already known to have desirable properties is kept fixed. A library of the complementary chain is inserted into the same vector. Suitable partners for the fixed chain are selected following display on the surface of bacteriophage.

[0037] In a fifth approach (see example 38), to improve the chances of recovering original pairs, the complexity of the combinatorial libraries can be reduced by using small B populations of B-lymphocytes selected for binding to a desired antigen. The cells provide e.g. mRNA or DNA, for preparing libraries of antibody genes for display on phage. This technique can be used in combination with the above mentioned four approaches for selection of antibody specificities.

[0038] Phagemids have been mentioned above. The applicants have realised and demonstrated that in many cases phagemids will be preferred to phage for cloning antibodies because it is easier to use them to generate more com-

prehensive libraries of the immune repertoire. This is because the phagemid DNA is approximately 100 times more efficient than bacteriophage DNA in transforming bacteria (see example 15). Also, the use of phagemids gives the ability to vary the number of gene III binding molecule fusion proteins displayed on the surface of the bacteriophage (see example 13). For example, in a system comprising a bacterial cell containing a phagemid encoding a gene III fusion protein and infected with a helper phage, induction of expression of the gene III fusion protein to different extents, will determine the number of gene III fusion proteins present in the space defined between the inner and outer bacterial membranes following superinfection. This will determine the ratio of gene III fusion protein to native gene III protein displayed by the assembled phage.

[0039] Expressing a single fusion protein per virion may aid selection of antibody specificities on the basis of affinity by avoiding the 'avidity' effect where a phage expressing two copies of a low affinity antibody would have the same apparent affinity as a phage expressing one copy of a higher affinity antibody. In some cases however, it will be important to display all the gene III molecules derived by superinfection of cells containing phagemids to have fusions (e.g. for selecting low affinity binding molecules or improving sensitivity on ELISA). One way to do this is to superinfect with a bacteriophage which contains a defective gene III. The applicants have therefore developed and used a phage which is deleted in gene III. This is completely novel.

[0040] The demonstration that a functional antigen-binding domain can be displayed on the surface of phage, has implications beyond the construction of novel antibodies. For example, if other protein domains can be displayed at the surface of a phage, phage vectors could be used to clone and select genes by the binding properties of the displayed protein. Furthermore, variants of proteins, including epitope libraries built into the surface of the protein, could be made and readily selected for binding activities. In effect, other protein architectures might serve as "nouvelle" antibodies.

[0041] The technique provides the possibility of building antibodies from first principles, taking advantage of the structural framework on which the antigen binding loops fold. In general, these loops have a limited number of conformations which generate a variety of binding sites by alternative loop combinations and by diverse side chains. Recent successes in modelling antigen binding sites augurs well for *de novo* design. In any case, a high resolution structure of the antigen is needed. However, the approach is attractive for making e.g. catalytic antibodies, particularly for small substrates. Here side chains or binding sites for prosthetic groups might be introduced, not only to bind selectively to the transition state of the substrate, but also to participate directly in bond making and breaking. The only question is whether the antibody architecture, specialised for binding, is the best starting point for building catalysts. Genuine enzyme architectures, such as the triose phosphate isomerase (TIM) barrel, might be more suitable. Like antibodies, TIM enzymes also have a framework structure (a barrel of β -strands and α -helices) and loops to bind substrate. Many enzymes with a diversity of catalytic properties are based on this architecture and the loops might be manipulated independently on the frameworks for design of new catalytic and binding properties. The phage selection system as provided by the present disclosure can be used to select for antigen binding activities and the CDR loops thus selected, used on either an antibody framework or a TIM barrel framework. Loops placed on a e.g. a TIM barrel framework could be further modified by mutagenesis and subjected to further selection. Thus, there is no need to select for high affinity binding activities in a single step. The strategy of the immune system, in which low affinity evolves to high affinity seems more realistic and can be mimicked using this invention.

[0042] Although throughout this application, the applicants discuss the possibility of screening for higher affinity variants of pAbs, they recognise that in some applications, for example low affinity chromatography (Ohlson, S. et al Anal. Biochem. 169, p204-208 (1988)), it may be desirable to isolate lower affinity variants.

[0043] Examples 17 and 19 show that the present invention provides a way of producing antibodies with low affinities (as seen in the primary immune response or in unimmunised animals). This is made possible by displaying multiple copies of the antibody on the phage surface in association with gene III protein. Thus, pAbs allow genes for these antibodies to be isolated and if necessary, mutated to provide improved antibodies.

[0044] pAbs also allow the selection of antibodies for improved stability. It has been noted for many antibodies, that yield and stability are improved when the antibodies are expressed at 30°C rather than 37°C. If pAbs are displayed at 37°C, only those which are stable will be available for affinity selection. When antibodies are to be used *in vivo* for therapeutic or diagnostic purposes, increased stability would extend the half-life of antibodies in circulation.

[0045] It will often be necessary to increase the diversity of a population of genes cloned for the display of their proteins on phage or to mutate an individual nucleotide sequence. Although *in vitro* or *in vivo* mutagenesis techniques could be used for either purpose, a particularly suitable method would be to use mutator strains. A mutator strain is a strain which contains a genetic defect which causes DNA replicated within it to be mutated with respect to its parent DNA. Hence if a population of genes as gene III fusions is introduced into these strains it will be further diversified and can then be transferred to a non-mutator strain, if desired, for display and selection. Example 29 covers the use of mutator strains with phage antibodies (an example of *in vitro* mutagenesis and selection of phage antibodies is given in example 35).

Targeted gene transfer

[0046] A useful and novel set of applications makes use of the binding protein on the phage to target the phage genome to a particular cell or group of cells. For example, a pAb specific for a cell surface molecule could be used to bind to the target cell via the surface molecule. The phage could then be internalised, either through the action of the receptor itself or as the result of another event (e.g. an electrical discharge such as in the technique of electroporation). The phage genome would then be expressed if the relevant control signals (for transcription and translation and possibly replication) were present. This would be particularly useful if the phage genome contained a sequence whose expression was desired in the target cell (along with the appropriate expression control sequences). A useful sequence might confer antibiotic resistance to the recipient cell or label the cell by the expression of its product (e.g. if the sequence expressed a detectable gene product such as a luciferase, see White, M. et al, Techniques 2(4), p194-201 (1990)), or confer a particular property on the target cell (e.g. if the target cell was a tumour cell and the new sequence directed the expression of a tumour suppressing gene), or express an antisense construct designed to turn off a gene or set of genes in the target cell, or a gene or gene product designed to be toxic to the target cell.

[0047] Alternatively, the sequence whose expression is desired in the target cell can be encoded on a phagemid. The phagemid DNA may then be incorporated into a phage displaying an antibody specific for a cell surface receptor. For example, incorporation may be by superinfection of bacteria containing the phagemid, with a helper phage whose genome encodes the antibody fragment specific for the target cell. The package is then used to direct the phagemid to the target cell.

[0048] This technique of "targeted gene transfer" has a number of uses in research and also in therapy and diagnostics. For example, gene therapy often aims to target the replacement gene to a specific cell type that is deficient in its activity. Targeting pAbs provide a means of achieving this.

[0049] In diagnostics, phage specific for particular bacteria or groups of bacteria have been used to target marker genes, e.g. luciferase, to the bacterial host (see, for example, Ulitzer, S., and Kuhn, J., EPA 85303913.9). If the host range of the phage is appropriate, only those bacteria that are being tested for, will be infected by the phage, express the luciferase gene and be detected by the light they emit. This system has been used to detect the presence of Salmonella. One major problem with this approach is the initial isolation of a bacteriophage with the correct host range and then the cloning of a luciferase gene cassette into that phage, such that it is functional. The pAb system allows the luciferase cassette to be cloned into a well characterised system (filamentous phage) and allows simple selection of an appropriate host range, by modifying the antibody (or other binding molecule) specificity that the pAb encodes.

[0050] The present applicants have also been able to develop novel selection systems and assay formats which depend on the unique properties of these pAbs.

TERMINOLOGY

[0051] Much of the terminology discussed in this section has been mentioned in the text where appropriate.

Specific Binding Pair

[0052] This describes a pair of molecules (each being a member of a specific binding pair) which are naturally derived or synthetically produced. One of the pair of molecules, has an area on its surface, or a cavity which specifically binds to, and is therefore defined as complementary with a particular spatial and polar organisation of the other molecule, so that the pair have the property of binding specifically to each other. Examples of types of specific binding pairs are antigen-antibody, biotin-avidin, hormone-hormone receptor, receptor-ligand, enzyme-substrate, IgG-protein A.

Multimeric Member

[0053] This describes a first polypeptide which will associate with at least a second polypeptide, when the polypeptides are expressed in free form and/or on the surface of a substrate. The substrate may be provided by a bacteriophage. Where there are two associated polypeptides, the associated polypeptide complex is a dimer, where there are three, a trimer etc. The dimer, trimer, multimer etc or the multimeric member may comprise a member of a specific binding pair.

[0054] Example multimeric members are heavy domains based on an immunoglobulin molecule, light domains based on an immunoglobulin molecule, T-cell receptor subunits.

Package

[0055] This describes a filamentous bacteriophage particle in which the particle is displaying a member of a specific binding pair at its surface. The package is a bacteriophage which displays an antigen binding domain at its surface.

This type of package has been called a phage antibody (pAb).

Antibody

- 5 **[0056]** This describes an immunoglobulin whether natural or partly or wholly synthetically produced. The term also covers any protein having a binding domain which is homologous to an immunoglobulin binding domain. These proteins can be derived from natural sources, or partly or wholly synthetically produced.

[0057] Example antibodies are the immunoglobulin isotypes and the Fab, F(ab')₂, scFv, Fv, dAb, Fd fragments.

10 Immunoglobulin Superfamily

[0058] This describes a family of polypeptides, the members of which have at least one domain with a structure related to that of the variable or constant domain of immunoglobulin molecules. The domain contains two β-sheets and usually a conserved disulphide bond (see A.F. Williams and A.N. Barclay 1988 Ann. Rev Immunol. 6 381-405).

- 15 **[0059]** Example members of an immunoglobulin superfamily are CD4, platelet derived growth factor receptor (PDGFR), intercellular adhesion molecule. (ICAM). Except where the context otherwise dictates, reference to immunoglobulins and immunoglobulin homologs in this application includes members of the immunoglobulin superfamily and homologs thereof.

20 Homologs

[0060] This term indicates polypeptides having the same or conserved residues at a corresponding position in their primary, secondary or tertiary structure. The term also extends to two or more nucleotide sequences encoding the homologous polypeptides.

- 25 **[0061]** Example homologous peptides are the immunoglobulin isotypes.

Functional

- 30 **[0062]** In relation to a sbp member displayed on the surface of a filamentous bacteriophage particle, means that the sbp member is presented in a folded form in which its specific binding domain for its complementary sbp member is the same or closely analogous to its native configuration, whereby it exhibits similar specificity with respect to the complementary sbp member. In this respect, it differs from the peptides of Smith et al, supra, which do not have a definite folded configuration and can assume a variety of configurations determined by the complementary members with which they may be contacted.

35 Genetically diverse population

[0063] In connection with sbp members or polypeptide components thereof, this is referring not only to diversity that can exist in the natural population of cells or organisms, but also diversity that can be created by artificial mutation in vitro or in vivo.

- 40 **[0064]** Mutation in vitro may for example, involve random mutagenesis using oligonucleotides having random mutations of the sequence desired to be varied. In vivo mutagenesis may for example, use mutator strains of host micro-organisms to harbour the DNA (see Example 38 below).

45 Domain

[0065] A domain is a part of a protein that is folded within itself and independently of other parts of the same protein and independently, of a complementary binding member.

50 Folded Unit

[0066] This is a specific combination of an α-helix and/or β-strand and/or β-turn structure. Domains and folded units contain structures that bring together amino acids that are not adjacent in the primary structure.

55 Free Form

[0067] This describes the state of a polypeptide which is not displayed by a replicable genetic display package.

Conditionally Defective

[0068] This describes a gene which does not express a particular polypeptide under one set of conditions, but expresses it under another set of conditions. An example, is a gene containing an amber mutation expressed in non-suppressing or suppressing hosts respectively.

[0069] Alternatively, a gene may express a protein which is defective under one set of conditions, but not under another set. An example is a gene with a temperature sensitive mutation.

Suppressible Translational Stop Codon

[0070] This describes a codon which allows the translation of nucleotide sequences downstream of the codon under one set of conditions, but under another set of conditions translation ends at the codon. Example of suppressible translational stop codons are the amber, ochre and opal codons.

Mutator Strain

[0071] This is a host cell which has a genetic defect which causes DNA replicated within it to be mutated with respect to its parent DNA. Example mutator strains are NR9046mutD5 and NR9046 mut T1 (see Example 38).

Helper Phage

[0072] This is a phage which is used to infect cells containing a defective phage genome and which functions to complement the defect. The defective phage genome can be a phagemid or a phage with some function encoding gene sequences removed. Examples of helper phages are M13K07, M13K07 gene III no. 3; and phage displaying or encoding a binding molecule fused to a capsid protein.

Vector

[0073] This is a DNA molecule, capable of replication in a host organism, into which a gene is inserted to construct a recombinant DNA molecule.

Phage Vector

[0074] This is a vector derived by modification of a phage genome, containing an origin of replication for a bacteriophage, but not one for a plasmid.

Phagemid Vector

[0075] This is a vector derived by modification of a plasmid genome, containing an origin of replication for a bacteriophage as well as the plasmid origin of replication.

Secreted

[0076] This describes a bacteriophage particle or molecule that associates with the member of a sbp displayed on the bacteriophage particle in which the sbp member and/or the molecule, have been folded and the package assembled externally to the cellular cytosol.

Repertoire of Rearranged Immunoglobulin Genes

[0077] A collection of naturally occurring nucleotides eg DNA sequences which encoded expressed immunoglobulin genes in an animal. The sequences are generated by the *in vivo* rearrangement of eg V, D and J segments for H chains and eg the V and J segments for L chains. Alternatively the sequences may be generated from a cell line immunised *in vitro* and in which the rearrangement in response to immunisation occurs intracellularly.

Library

[0078] A collection of nucleotide eg DNA, sequences within clones.

Repertoire of Artificially Rearranged Immunoglobulin Genes

[0079] A collection of nucleotide eg DNA, sequences derived wholly or partly from a source other than the rearranged immunoglobulin sequences from an animal. This may include for example, DNA sequences encoding VH domains by combining unrearranged V segments with D and J segments and DNA sequences encoding VL domains by combining V and J segments.

[0080] Part or all of the DNA sequences may be derived by oligonucleotide synthesis.

Secretory Leader Peptide

[0081] This is a sequence of amino acids joined to the N-terminal end of a polypeptide and which directs movement of the polypeptide out of the cytosol.

Eluant

[0082] This is a solution used to breakdown the linkage between two molecules. The linkage can be a non-covalent or covalent bond(s). The two molecules can be members of a sbp.

Derivative

[0083] This is a substance which derived from a polypeptide which is encoded by the DNA within a selected bacteriophage particle. The derivative polypeptide may differ from the encoded polypeptide by the addition, deletion, substitution or insertion of amino acids, or by the linkage of other molecules to the encoded polypeptide. These changes may be made at the nucleotide or protein level. For example the encoded polypeptide may be a Fab fragment which is then linked to an Fc tail from another source. Alternatively markers such as enzymes, flouresceins etc may be linked to eg Fab, scFv fragments.

[0084] The present invention provides a method for producing a filamentous bacteriophage particle displaying at its surface a binding molecule specific for a particular target epitope or antigen, which method comprises the steps of:

producing a population of filamentous bacteriophage particles displaying at their surface a population of binding molecules having a range of binding specificities, wherein the binding molecules are Fab antibody molecules able to bind target epitope or antigen, and wherein each filamentous bacteriophage particle contains a phagemid genome comprising nucleic acid with a nucleotide sequence encoding the binding molecule expressed from the nucleic acid and displayed by the particle at its surface;

selecting for a filamentous bacteriophage particle displaying a binding molecule with a desired specificity by contacting the population of filamentous bacteriophage particles with a target epitope or antigen so that individual binding molecules displayed on filamentous bacteriophage particles with the desired specificity bind to said target epitope or antigen.

[0085] The method may comprise one or more of the additional steps of (i) separating any bound filamentous bacteriophage particles from the target epitope or antigen; and (ii) recovering any separated filamentous bacteriophage particles displaying a binding molecule with the desired specificity. The selection step may isolate the nucleotide sequence encoding the binding molecule of desired specificity, by virtue of said binding molecule being expressed in association with the surface of the filamentous bacteriophage particle in which said encoding nucleic acid is contained. Accordingly, the present invention also provides a method for producing a binding molecule specific for a particular target epitope or antigen, which method comprises:

performing a method as set out in the preceding paragraph and including the above steps (i) and (ii);
isolating from separated filamentous bacteriophage particles so recovered nucleic acid encoding the binding molecule;

inserting nucleic acid encoding the binding molecule, or a fragment or derivative thereof with binding specificity for the target epitope or antigen, in a recombinant system; and producing the binding molecule, or fragment or derivative thereof with binding specificity for the target epitope or antigen, in the recombinant system separate from filamentous bacteriophage particles.

[0086] A helper phage, or a plasmid expressing complementing phage genes, is used to help package said phagemid genome.

[0087] A library or genetically diverse population may be obtained from:

- (i) the repertoire of rearranged immunoglobulin genes of an animal immunised with complementary sbp member,
- (ii) the repertoire of rearranged immunoglobulin genes of an animal not immunised with complementary sbp member,
- (iii) a repertoire of artificially rearranged immunoglobulin gene or genes
- (iv) a repertoire of immunoglobulin homolog gene or genes; or
- (v) a mixture of any of (i), (ii), (iii) and (iv). The capsid protein may be absent, defective or conditionally defective in the helper phage.

[0088] The host cell may be a mutator strain which introduces genetic diversity into the sbp member nucleic acid.

[0089] The host may be a bacterium, and said component of the filamentous bacteriophage particle a capsid protein for the bacteriophage. The phage may be selected from the class I phages fd, M13, f1, If1, Ike, ZJ/Z, Ff and the class II phages Xf, Pf1 and Pf3. The phage may be fd or a derivative of fd. The derivative may be tetracycline resistant. The said sbp member or polypeptide chain thereof may be expressed as a fusion with the gene III capsid protein of phage fd or its counterpart in another filamentous phage. The sbp member or polypeptide chain thereof may be inserted in the N-terminal region of the mature capsid protein downstream of a secretory leader peptide. The sequence may be inserted after amino acid +1 of the mature protein. The site for insertion may be flanked by short sequences corresponding to sequences which occur at each end of the nucleic acid to be inserted. For example where the protein domain is an immunoglobulin domain, the insertion site in the phage may be flanked by nucleotide sequences which code for the first five amino acids and the last five amino acids of the Ig domain. Such flanking nucleotide sequences are shown in figure 4(2) B and C, wherein the site-flanking nucleotide sequences encode amino acid sequences QVQLQ and VTVSS which occur at either end of the VH domain, or QVQLQ and LEIKR which occur at either end of the Fv (combined VH + VL) domain. Each of these sequences flanking the insertion site may include a suitable cleavage site, as shown in Fig 4.

[0090] Alternatively, the flanking nucleotide sequences shown in figure 4(2)B and C as described above, may be used to flank the insertion site for any nucleic acid to be inserted, whether or not that nucleic acid codes an immunoglobulin.

[0091] The host may be E.coli.

[0092] Nucleic acid encoding an sbp member polypeptide may be linked downstream to a viral capsid protein through a suppressible translational stop codon.

[0093] In a method for producing a binding molecule as defined above, the gene sequence encoding the binding molecule of desired specificity is separated from a general population of filamentous bacteriophage particle having a range of specificities, by the fact of its binding to a specific target (eg the antigen or epitope). Thus the filamentous bacteriophage particle formed by said expression may be selected or screened to provide an individual sbp member or a selected mixed population of said sbp members associated in their respective particles with nucleic acid encoding said sbp member or a polypeptide chain thereof. The particles may be selected by affinity with a member complementary to said sbp member.

[0094] Any particles bound to said second member may be recovered by washing with an eluant. The washing conditions may be varied in order to obtain particles with different binding affinities for said epitope. Alternatively, to obtain eg high affinity particles the complementary member (eg an epitope) may be presented to the population of particles (pAbs) already bound to a binding member in which case pAbs with a higher affinity for the epitope will displace the already bound binding member. Thus the eluant may contain a molecule which competes with said particles for binding to the complementary sbp member. The particles may be applied to said complementary sbp member in the presence of a molecule which competes with said package for binding to said complementary sbp member. Nucleic acid derived from a selected or screened particle may be used to express said sbp member or a fragment or derivative thereof in a recombinant host organism. Nucleic acid from one or more particles may be taken and used to provide encoding nucleic acid in a further said method to obtain an individual sbp member or a mixed population of sbp members, or encoding nucleic acid therefor. The expression end product may be modified to produce a derivative thereof.

[0095] The expression end product or derivative thereof may be used to prepare a therapeutic or prophylactic medicament or a diagnostic product.

[0096] The present invention may employ a helper phage whose genome lacks nucleic acid encoding one of its capsid proteins, or whose encoding nucleic acid therefor is conditionally defective, or which encodes said capsid protein in defective or conditionally defective form.

[0097] The present invention may employ recombinant E.coli TG1 M13K07 gIII No. 3 (NCTC 12478).

[0098] The present invention also provides a filamentous bacteriophage particle displaying on its surface a binding molecule which is a Fab or single-chain Fv antibody molecule comprising a binding domain able to bind target epitope or antigen, the particle containing a phagemid genome comprising nucleic acid with a nucleotide sequence encoding the binding molecule expressed from the nucleic acid and displayed by the particle at its surface.

[0099] In the above methods, the binding molecule may be either naturally derived or synthetic or a combination of

both.

[0100] The present invention provides filamentous bacteriophage particles as defined above, and a population of such filamentous bacteriophage particles displaying a population of said binding molecules having a range of binding specificities. The present invention also provides a method for producing Fab antibody molecules, fragments and derivatives thereof. The derivatives may comprise members of the specific binding pairs fused to another molecule such as an enzyme or a Fc tail.

[0101] The invention may employ kits for carrying out the methods hereof. The kits will include the necessary vectors. One such vector will typically have an origin of replication for single stranded bacteriophage and either contain the sbp member nucleic acid or have a restriction site for its insertion in the 5' end region of the mature coding sequence of a phage capsid protein, and with a secretory leader coding sequence upstream of said site which directs a fusion of the capsid protein exogenous polypeptide to the periplasmic space.

[0102] The restriction sites in the vectors are preferably those of enzymes which cut only rarely in protein coding sequences.

[0103] The kit preferably includes a phagemid vector which may have the above characteristics, or may contain, or have a site for insertion, of sbp member nucleic acid for expression of the encoded polypeptide in free form.

[0104] The kits will also contain ancillary components required for carrying out the method, the nature of such components depending of course on the particular method employed.

[0105] Useful ancillary components may comprise helper phage, PCR primers, and buffers and enzymes of various kinds.

[0106] PCR primers and associated reagents for use where the sbp members are antibodies may have the following characteristics:

(i) primers having homology to the 5' end of the sense or anti-sense strand of sequences encoding domains of antibodies; and

(ii) primers including tag sequences 5' to these homologous sequences which, incorporate restriction sites to allow insertion into vectors; together with sequences to allow assembly of amplified VH and VL regions to enable expression as Fv, scFv or Fab fragments.

[0107] Buffers and enzymes are typically used to enable preparation of nucleotide sequences encoding Fv, scFv or Fab fragments derived from rearranged or unrearranged immunoglobulin genes according to the strategies described herein.

[0108] The applicants have chosen the filamentous F-specific bacteriophages as an example of the type of phage which could provide a vehicle for the display of binding molecules e.g. antibodies and antibody fragments and derivatives thereof, on their surface and facilitate subsequent selection and manipulation.

[0109] The F-specific phages (e.g. f1, fd and M13) have evolved a method of propagation which does not kill the host cell and they are used commonly as vehicles for recombinant DNA (Kornberg, A., DNA Replication, W.H. Freeman and Co., San Francisco, 1980). The single stranded DNA genome (approximately 6.4 Kb) of fd is extruded through the bacterial membrane where it sequesters capsid sub-units, to produce mature virions. These virions are 6 nm in diameter, 1µm in length and each contain approximately 2,800 molecules of the major coat protein encoded by viral gene VIII and four molecules of the adsorption molecule gene III protein (g3p) the latter is located at one end of the virion. The structure has been reviewed by Webster et al., 1978 in The Single Stranded DNA Phages, 557-569, Cold Spring Harbor Laboratory Press. The gene III product is involved in the binding of the phage to the bacterial F-pilus.

[0110] Although these phages do not kill their host during normal replication, disruption of some of their genes can lead to cell death (Kornberg, A., 1980 supra.) This places some restraint on their use. The applicants have recognized that gene III of phage fd is an attractive possibility for the insertion of biologically active foreign sequences. There are however, other candidate sites including for example gene VIII and gene VI.

[0111] The protein itself is only a minor component of the phage coat and disruption of the gene does not lead to cell death (Smith, G. 1988, Virology 167: 156-165). Furthermore, it is possible to insert some foreign sequences (with no biological function) into various positions within this gene (Smith, G. 1985 Science 228: 1315-1317., Parmley, S.F. and Smith, G.P. Gene: 73 (1988) p. 305-318., and de la Cruz, V.F., et al., 1988, J. Biol. Chem., 263: 4318-4322). Smith et al described the display of peptides on the outer surface of phage but they did not describe the display of protein domains. Peptides can adopt a range of structures which can be different when in free solution, than when bound to, for example, an antibody, or when forming part of a protein (Stanfield, R.I. et al., (1990) Science 248, p712-719). Proteins in general have a well defined tertiary structure and perform their biological function only when adopting this structure. For example, the structure of the antibody D1.3 has been solved in the free form and when bound to antigen (Bhat, T.N: et al., (1990) Nature 347, p483-485). The gross structure of the protein is identical in each instance with only minor variations around the binding site for the antigen. Other proteins have more substantial conformation changes on binding of ligand, for instance the enzymes hexokinase and pyruvate dehydrogenase during their catalytic cycle,

but they still retain their overall pattern of folding. This structural integrity is not confined to whole proteins, but is exhibited by protein domains. This leads to the concept of a folded unit which is part of a protein, often a domain, which has a well defined primary, secondary and tertiary structure and which retains the same overall folding pattern whether binding to a binding partner or not. The only gene sequence that Smith et al., described that was of sufficient size to encode a domain (a minimum of perhaps 50 amino acids) was a 335bp fragment of a β -galactosidase corresponding to nucleotides 861-1195 in the β -galactosidase gene sequence (Pamley, S. Smith, G. P. 1988 supra. This would encode 112 amino acids of a much larger 380 amino acid domain. Therefore, prior to the present application, no substantially complete domain or folded unit had been displayed on phage. In these cases, although the infectivity of the virion was disrupted, the inserted sequences could be detected on the phage surface by use of e.g. antibodies.

[0112] The protein encoded by gene III has several domains (Pratt, D., et al., 1969 *Virology* 39:42-53., Grant, R.A., et al., 1981, *J. Biol. Chem.* 256: 539-546 and Armstrong, J., et al., *FEES Lett.* 135: 167-172 1981.) including: (i) a signal sequence that directs the protein to the cell membrane and which is then cleaved off; (ii) a domain that anchors the mature protein into the bacterial cell membrane (and also the phage coat); and (iii) a domain that specifically binds to the phage receptor, the F-pilus of the host bacterium. Short sequences derived from protein molecules have been inserted into two places within the mature molecule (Smith, G., 1985 supra., and Pamley, S.F. and Smith G.P., 1988 supra.). Namely, into an inter-domain region and also between amino acids 2 and 3 at the N-terminus. The insertion sites at the N-terminus were more successful in maintaining the structural integrity of the gene III protein and displaying the peptides on the surface of the phage. By use of antisera specific for the peptides, the peptides inserted into this position were shown to be on the surface of the phage. These authors were also able to purify the phage, using this property. However, the peptides expressed by the phage, did not possess measurable biological functions of their own.

[0113] Retaining the biological function of a molecule when it is expressed in a radically different context to its natural state is difficult. The demands on the structure of the molecule are heavy. In contrast, retaining the ability to be bound by specific antisera is a passive process which imposes far less rigorous demands on the structure of the molecule. For example, it is the rule rather than the exception that polyclonal antisera will recognise totally denatured, and biologically inactive, proteins on Western blots (see for example, Harlow, E. and Lane, D., *Antibodies, a Laboratory Manual*, Cold Spring Harbor Laboratory Press 1988). Therefore, the insertion of peptides into a region that allows their structure to be probed with antisera teaches only that the region allows the inserted sequences to be exposed and does not teach that the region is suitable for the insertion of large sequences with demanding structural constraints for the display of a molecule with a biological or binding function. In particular, it does not teach that domains or folded units of proteins can be displayed from sequences inserted in this region.

[0114] This experience with Western blots is a graphic practical demonstration which shows that retaining the ability to be bound by specific antisera imposes far less rigorous demands on the structure of a polypeptide, than does folding for the retention of a biological function.

[0115] Studies have been carried out, in which *E.coli* have been manipulated to express the protein β -adrenergic receptor as a fusion with the outer membrane protein lamB. The β -adrenergic receptor was expressed in a functional form as determined by the presence of binding activity. However, when an equivalent antibody fusion was made with lamB, the antibody fusion was toxic to the host cell.

[0116] The applicants have investigated the possibility of inserting the gene coding sequence for biologically active antibody fragments into the gene III region of fd to express a large fusion protein. As is apparent from the previous discussion, this approach makes onerous demands on the functionality of the fusion protein. The insertion is large, encoding antibody fragments of at least 100-200 amino acids; the antibody derived domain must fold efficiently and correctly to display antigen-binding; and most of the functions of gene III must be retained. The applicants approach to the construction of the fusion molecule was designed to minimise the risk of disrupting these functions. In an embodiment of the invention, the initial vector used was fd-tet (Zacher, A.N., et al., 1980, *Gene* 9: 127-140) a tetracycline resistant version of fd bacteriophage that can be propagated as a plasmid that confers tetracycline resistance to the infected *E.coli* host. The applicants chose to insert after the signal sequence of the fd gene III protein for several reasons. In particular, the applicants chose to insert after amino acid 1 of the mature protein to retain the context for the signal peptidase cleavage. To retain the structure and function of gene III itself, the majority of the original amino acids are synthesized after the inserted immunoglobulin sequences. The inserted immunoglobulin sequences were designed to include residues from the switch region that links VH-VL to CH1-CL (Lesk, A., and Chothia, C., *Nature* 335, 188-190, 1988).

[0117] Surprisingly, by manipulating gene III of bacteriophage fd, the present applicants have been able to construct a bacteriophage that displays on its surface large biologically functional antibody, enzyme, and receptor molecules whilst remaining intact and infectious. Furthermore, the phages bearing antibodies of desired specificity, can be selected from a background of phages not showing this specificity.

[0118] The sequences coding for a population of antibody molecules and for insertion into the vector to give expression of antibody binding functions on the phage surface can be derived from a variety of sources. For example, immunised or non-immunised rodents or humans, and from organs such as spleen and peripheral blood lymphocytes. The

coding sequences are derived from these sources by techniques familiar to those skilled in the art (Orlandi, R., et al., 1989 supra; Larrick, J.W., et al., 1989 supra; Chiang, Y.L., et al., 1989 Bio Techniques 7, p. 360-366; Ward, E.S., et al., 1989 supra; Sastry, L., et al., 1989 supra.) or by novel linkage strategies described in examples 12, 26, 30 and 32. Novel strategies are described in examples 7, 21, 26 and 30 for displaying dimeric molecules eg Fab and Fv fragments on the surface of a phage. Each individual pAb in the resulting library of pAbs will express antibodies or antibody derived fragments that are monoclonal with respect to their antigen-binding characteristics.

[0119] The disclosure made by the present applicants is important and provides a significant breakthrough in the technology relating to the production of biological binding molecules, their fragments and derivatives by the use of recombinant methods.

[0120] In standard recombinant techniques for the production of antibodies, an expression vector containing sequences coding for the antibody polypeptide chains is used to transform e.g. E.coli. The antibody polypeptides are expressed and detected by use of standard screening systems. When the screen detects an antibody polypeptide of the desired specificity, one has to return to the particular transformed E.coli expressing the desired antibody polypeptide. Furthermore, the vector containing the coding sequence for the desired antibody polypeptide then has to be isolated for use from E.coli in further processing steps.

[0121] In the present invention however, the desired antibody polypeptide when expressed, is already packaged with its gene coding sequence. This means that when the an antibody polypeptide of desired specificity is selected, there is no need to return to the original culture for isolation of that sequence. Furthermore, in previous methods in standard recombinant techniques, each clone expressing antibody needs to be screened individually. The present application provides for the selection of clones expressing antibodies with desired properties and thus only requires screening of clones from an enriched pool.

[0122] Because a pAb is a novel structure that displays a member of a specific binding pair (an antibody of monoclonal antigen-binding specificity) at the surface of a relatively simple replicable structure also containing the genetic information encoding the member, pAbs that bind to the complementary member of the specific binding pair (eg antigen) can be recovered very efficiently by either eluting off the complementary member using for example diethylamine, high salt etc and infecting suitable bacteria, or by denaturing the structure, and specifically amplifying the sequences encoding the member using PCR. That is, there is no necessity to refer back to the original bacterial clone that gave rise to the pAb.

[0123] For some purposes, for example immunoprecipitation and some diagnostic tests, it is advantageous to use polyclonal antibodies or antibody fragments. The present invention allows this to be achieved by either selection of an enriched pool of pAbs with desired properties or by mixing individually isolated clones with desired properties. The antibodies or antibody fragments may then be expressed in soluble form if desired. Such a selected polyclonal pAb population can be grown from stocks of phage, bacteria containing phagemids or bacteria expressing soluble fragments derived from the selected polyclonal population. Thus a reagent equivalent to a polyclonal antiserum is created which can be replicated and routinely manufactured in culture without use of animals.

SELECTION FORMATS AND AFFINITY MATURATION

[0124] Individual pAbs expressing the desired specificity eg for an antigen, can be isolated from the complex library using the conventional screening techniques (e.g. as described in Harlow, E., and Lane, D., 1988, supra Gherardi, E et al. 1990. J. Immunol. meth. 126 p61-68).

[0125] The applicants have also devised a series of novel selection techniques that are practicable only because of the unique properties of filamentous bacteriophage particles. The general outline of some screening procedures is illustrated in figure 2.

[0126] The population/library of pAbs to be screened could be generated from immunised or other animals; or be created in vitro by. mutagenising pre-existing phage antibodies (using techniques well-known in the art such as oligonucleotide directed mutagenesis (Sambrook, J., et al., 1989 Molecular Cloning a Laboratory Manual, Cold Spring Harbor Laboratory Press). This population can be screened in one or more of the formats described below with reference to figure 2, to derive those individual pAbs whose antigen binding properties are different from sample c.

Binding Elution

[0127] Figure 2(i) shows antigen (ag) bound to a solid surface (s) the solid surface (s) may be provided by a petri dish, chromatography beads, magnetic beads and the like. The population/library of pAbs is then passed over the ag, and those individuals p that bind are retained after washing, and optionally detected with detection system d. A detection system based upon anti-fd antisera is illustrated in more detail below in example 4. If samples of bound population p are removed under increasingly stringent conditions, the binding affinity represented in each sample will increase. Conditions of increased stringency can be obtained, for example, by increasing the time of soaking or changing the

pH of the soak solution, etc.

Competition

5 **[0128]** Referring to figure 2(ii) antigen ag can be bound to a solid support s and bound to saturation by the original binding molecule c. If a population of mutant pAb (or a set of unrelated pAbs) is offered to the complex, only those that have higher affinity for antigen ag than c will bind. In most examples, only a minority of population c will be displaced by individuals from population p. If c is a traditional antibody molecule, all bound material can be recovered and bound p recovered by infecting suitable bacteria and/or by use of standard techniques such as PCR.

10 **[0129]** An advantageous application is where ag is an antibody and c its antigen. The recovered bound population p is then an anti-idiotypic antibody which have numerous uses in research and the diagnostic and pharmaceutical industries.

[0130] At present it is difficult to select directly for anti-idiotypic antibodies. pAbs would give the ability to do this directly by binding pAb libraries (eg a naive library) to B cells (which express antibodies on their surface) and isolating those phage that bound well.

[0131] In some instances it may prove advantageous to preselect population p. For example, in the anti-idiotypic example above, p can be absorbed against a related antibody that does not bind the antigen.

[0132] However, if c is a pAb, then either or both c and p can advantageously be marked in some way to both distinguish and select for bound p over bound c. This marking can be physical, for example, by pre-labelling p with biotin; or more advantageously, genetic. For example, c can be marked with an EcoB restriction site, whilst p can be marked with an EcoK restriction site (see Carter, P. et al., 1985, Nucl. Acids Res. 13, 4431-4443). When bound p-c are eluted from the antigen and used to infect suitable bacteria, there is restriction (and thus no growth) of population c (i.e. EcoB restricting bacteria in this example). Any phage that grew, would be greatly enriched for those individuals from p with higher binding affinities. Alternatively, the genetic marking can be achieved by marking p with new sequences, which can be used to specifically amplify p from the mixture using PCR.

25 **[0133]** Since the bound pAbs can be amplified using for example PCR or bacterial infection, it is also possible to rescue the desired specificity even when insufficient individuals are bound to allow detection via conventional techniques.

[0134] The preferred method for selection of a phage displaying a protein molecule with a desired specificity or affinity will often be elution from an affinity matrix with a ligand (eg example 17). Elution with increasing concentrations of ligand should elute phage displaying binding molecules of increasing affinity. However, when eg a pAb binds to its antigen with high affinity or avidity (or another protein to its binding partner) it may not be possible to elute the pAb from an affinity matrix with molecule related to the antigen. Alternatively, there may be no suitable specific eluting molecule that can be prepared in sufficiently high concentration. In these cases it is necessary to use an elution method which is not specific to eg the antigen-antibody complex. Some of the non-specific elution methods generally used reduce phage viability for instance, phage viability is reduced with time at pH12 (Rossomando, E.F. and Zinder N.D. J. Mol.Biol. 36 387-399 1968). There may be interactions between eg antibodies and affinity matrices which cannot be disrupted without completely removing phage infectivity. In these cases a method is required to elute phage which does not rely on disruption of eg the antibody - antigen interaction. A method was therefore devised which allows elution of bound pAbs under mild conditions (reduction of a dithiol group with dithiothreitol) which do not disrupt phage structure (example 37).

[0135] This elution procedure is just one example of an elution procedure under mild conditions. A particularly advantageous method would be to introduce a nucleotide sequence encoding amino acids constituting a recognition site for cleavage by a highly specific protease between the foreign gene inserted, in this instance a gene for an antibody fragment, and the sequence of the remainder of gene III. Examples of such highly specific proteases are Factor X and thrombin. After binding of the phage to an affinity matrix and elution to remove non-specific binding phage and weak binding phage, the strongly bound phage would be removed by washing the column with protease under conditions suitable for digestion at the cleavage site. This would cleave the antibody fragment from the phage particle eluting the phage. These phage would be expected to be infective, since the only protease site should be the one specifically introduced. Strongly binding phage could then be recovered by infecting eg. E.coli TG1 cells.

50 **[0136]** An alternative procedure to the above is to take the affinity matrix which has retained the strongly bound pAb and extract the DNA, for example by boiling in SDS solution. Extracted DNA can then be used to directly transform E. coli host cells or alternatively the antibody encoding sequences can be amplified, for example using PCR with suitable primers such as those disclosed herein, and then inserted into a vector for expression as a soluble antibody for further study or a pAb for further rounds of selection.

[0137] Another preferred method for selection according to affinity would be by binding to an affinity matrix containing low amounts of ligand.

[0138] If one wishes to select from a population of phages displaying a protein molecule with a high affinity for its

ligand, a preferred strategy is to bind a population of phage to an affinity matrix which contains a low amount of ligand. There is competition between phage, displaying high affinity and low affinity proteins, for binding to the ligand on the matrix. Phage displaying high affinity protein is preferentially bound and low affinity protein is washed away. The high affinity protein is then recovered by elution with the ligand or by other procedures which elute the phage from the affinity matrix (example 35 demonstrates this procedure).

[0139] In summary then, for recovery of the packaged DNA from the affinity step, the package can be simply eluted, it can be eluted in the presence of a homologous sbp member which competes with said package for binding to a complementary sbp member; it could be removed by boiling, it could be removed by proteolytic cleavage of the protein; and other methods will be apparent to those skilled in the art eg. destroying the link between the substrate and complementary sbp member to release said packaged DNA and sbp member. At any rate, the objective is to obtain the DNA from the package so that it can be used directly or indirectly, to express the sbp member encoded thereby.

[0140] The efficiency of this selection procedure for pAbs and the ability to create very large libraries means that the immunisation techniques developed to increase the proportion of screened cells producing antibodies of interest will not be an absolute requirement. The technique allows the rapid isolation of binding specificities eg antigen-binding specificities, including those that would be difficult or even unobtainable by conventional techniques, for example, catalytic or anti-idiotypic antibodies. Removal of the animal altogether is now possible, once a complete library-of the immune repertoire has been constructed.

The novel structure of the pAb molecule can be used in a number of other applications, some examples of which are:

Signal Amplification

[0141] Acting as a novel molecular entity in itself pAbs combine the ability to bind a specific molecule eg antigen with amplification, if the major coat protein is used to attach another moiety. This moiety can be attached via immunological, chemical, or any other means and can be used, for example, to label the complex with detection reagents or cytotoxic molecules for use in vivo or in vitro.

Physical Detection

[0142] The size of the pAbs can be used as a marker particularly with respect to physical methods of detection such as electron microscopy and/or some biosensors, e.g. surface plasmon resonance.

Diagnostic Assays

[0143] The pAbs also have advantageous uses in diagnostic assays, particularly where separation can be effected using their physical properties for example centrifugation, filtration etc.

[0144] In order that the invention is more fully understood, embodiments will now be described in more detail by way of example only and not by way of limitation with reference to the figures described below.

[0145] In all embodiments of the invention filamentous bacteriophage particles containing a phagemid genome are employed. Display at the surface of particles containing a phage genome is not part of the invention and mention of this in this document, including the experimental examples, is for the purpose of illustration.

[0146] Figure 1 shows the basic structure of the simplest antibody molecule IgG.

[0147] Figure 2 shows schematically selection techniques which utilise the unique properties of pAbs; 2i) shows a binding/elution system; and (2ii) shows a competition system (p=pAb; ag=antigen to which binding by pAb is required; c=competitor population e.g. antibody, pAb, ligand; s=substrate (e.g. plastic beads etc); d=detection system).

[0148] Figure 3 shows the vector fd-tet and a scheme for the construction of vectors, fdTPs/Bs (for insertion of VH coding sequences) and fdTPs/Xh for the insertion of scFv coding sequences.

[0149] Figure 4 shows the nucleotide sequences for the oligonucleotides and vectors. All sequences are drawn 5' to 3' and are numbered according to Beck et al., 1978, Nucl. Acid Res., 5: 4495-4503. 4.1 shows the sequences of the oligonucleotides used for mutagenesis (oligo's 1 and 2) or sequencing (oligo 3). The sequences shown were synthesized on an Applied Biosystems, oligonucleotide synthesizer and are complementary to the single stranded form of fd-tet (they are in the anti-sense form with respect to gene III). 4.2 shows the sequences of the various constructs around the gene III insertion site. These sequences are drawn in the sense orientation with respect to gene III; (A) fd-tet (and fdTδBst) (B) fdTPs/Bs and (C) fdTPs/Xh. The key restriction enzyme sites are shown along with the immunoglobulin amino acids contributed by the vectors, (amino acid single letter code is used, see Harlow, E., and Lane, D., 1988 supra.).

[0150] Figure 5 shows the nucleotide and amino acid sequences for scFv in the vector scFvD1.3 myc. This gives the sequence of the anti-lysozyme single chain Fv and surrounding sequences in scFvD1.3 myc, showing the N-terminal pel B signal peptide sequence and the C-terminal myc tag sequence (Ward, E.S., et al., 1989, supra.). Also

shown is the peptide sequence linking the VH and VL regions. The amino acid sequence is represented above the nucleotide sequence by the single letter code, see Harlow, E., and Lane D., 1988 supra.

[0151] Figure 6 shows the binding of pAbs to lysozyme and the effect of varying the amount of supernatant. Each point is the average of duplicate samples. Lysozyme was coated at 1 mg/ml in 50 mM NaHCO₃.

[0152] Figure 7 shows the effect of varying the coating concentration of lysozyme or bovine serum albumin on the binding of pAbs to lysozyme in graphical form. Each point is the average of duplicate samples.

[0153] Figure 8 shows the sequence around the cloning site in gene III of fd-CAT2. Restriction enzyme sites are shown as well as the amino acids encoded by antibody derived sequences. These are flanked at the 5' end by the gene III signal peptide and at the 3' end by 3 alanine residues (encoded by the Not I restriction site) and the remainder of the mature gene III protein. The arrow shows the cleavage site for cutting of the signal peptide.

[0154] Figure 9 shows the binding of pAb (1.3) to lysozymes. Binding of phage as detected by ELISA to (a) hen egg-white lysozyme (HEL) (b) turkey egg-white lysozyme (TEL), (c) human lysozyme (HUL), (d) bovine serum albumin (BSA). A further control of (e) fdTPs/Bs to HEL.

[0155] Figure 10 shows a map of FabD1.3 in pUC19.

[0156] Figure 11 shows the ELISA results providing a comparison of lysozyme-binding by phage-Fab and phage-scFv. Vector=fdCAT2 (example 5); fdscFv(OX)=pAbNQ11 (Example 9); fdVHCHI (D1.3)=grown in normal cells (i.e. no L chain, see example 7); fdFab(D1.3) i.e. fdVHCHI (D1.3) grown in cells containing D1.3 L chain; fdscFv (D1.3)=pAbD1.3.

[0157] Figure 12 shows oligonucleotide probing of affinity purified phage. 10¹² phage in the ratio of 1 pAb (D1.3) in 4 x 10⁴ fdTPS/Bs phages were affinity purified and probed with an oligonucleotide specific for pAb (D1.3) A is a filter after one round of affinity purification (900 colonies total) and B is a filter after two rounds (372 colonies total).

[0158] Figure 13 shows the sequence of the anti-oxazolone antibody fragment NQ11 scFv. The sequence contributed by the linker is shown in the lower case. The sequence for VH is before the linker sequence and the sequence for VL is after the linker.

[0159] Figure 14 shows the ELISA results for binding pAb NQ11 and pAb D1.3 and vector fdTPs/xh to specified antigens.

[0160] Figure 15 shows the sequence surrounding the phoA insertion in fd-phoAla166. The restriction sites used for cloning are shown, as well as the amino acids encoded by phoA around the insertion site. The first five amino acids of the mature fusion come from gene III.

[0161] Figure 16(1) shows the structure of gene III and the native BamHI site into which a scFv coding sequence was inserted in example 11 and figure 16(2) shows the natural peptide linker sites A and B for possible insertion of scFv coding sequences.

[0162] Figure 17 shows schematically the protocol for PCR assembly of mouse VH and VLK repertoires for phage display described in example 12.

[0163] Figure 18 shows examples of the final products obtained with the procedure of example 12. Lanes a and b show the products of the initial PCR using heavy and light chain primers respectively; lane c shows the complete assembled 700bp product before final digestion with NotI and ApaI; M1, M2 markers Φ 174 Hae III digest and 123 base pair ladder (BRL Limited, P.O. Box 35, Washington Road, Paisley, Scotland) respectively.

[0164] Figure 19 shows the results of an ELISA of lysozyme binding by pCAT-3 scFv D1.3 phagemid in comparison with pCAT-3 vector (both rescued by M13K07) and fdCAT2 scFv D1.3 as described in example 13. The ELISA was performed as described in example 6 with modifications detailed in example 14.

[0165] Figure 20 shows the digestion pattern seen when individual clones, selected at random from a library of single chain Fv antibody genes derived from an immunised mouse; are digested with BstNI.

[0166] Figure 21 shows VH and VK gene sequences derived from the combinatorial library in example 17 and the hierarchical library in example 18.

[0167] Figure 22 shows a matrix of ELISA signals for clones derived from random combinatorial library. Designation of the clones is as in figure 21. The number of clones found with each combination is shown by the numerals.

[0168] Figure 23 shows a) the phagemid pHEN1 a derivative of pUC119 described in example 20; and b) the cloning sites in the phagemid pHEN.

[0169] Figure 24 The antibody constructs cloned into fd-CAT2 and pHEN1 for display on the surface of phage. Constructs I, II, III and IV were cloned into both fd-CAT2 (as ApaLI-NotI fragments) and pHEN1 (as SfiI-NotI fragments) and pHEN1 (as SfiI-NotI fragments). All the constructs contained the heavy chain (VH) and light chain (VK) variable regions of the mouse anti-phOx antibody NQ10.12.5. The constant domains were human CK and CH1 (γ 1 isotype).

[0170] Figure 25. Three ways of displaying antibody fragments on the surface of phage by fusion to gene III protein.

[0171] Figure 26. Western blot of supernatant taken from pHEN1-II(+) or pHEN1(-) cultures in E.coli HB2151, showing secretion of Fab fragment from pHEN1-II only. The anti-human Fab detects both H and L chain. Due to the attached c-myc tag, the L chain, highlighted by both anti-c-myc tag and anti-human CK antisera, is slightly larger (calculated Mr 24625) than the H chain (calculated Mr 23145).

[0172] Figure 27 is a plot showing the effect of lysozyme dilution on ratio of ELISA signals obtained using pAbD1.3 or soluble scFv D1.3.

[0173] Figure 28 is a plot showing the effect of lysozyme dilution on ELISA signals obtained using fdTscFvD1.3 and soluble scFvD1.3.

5 **[0174]** Figure 29 is a plot showing positive results from an ELISA screen of phage displaying scFv fragments derived from the cell line 013 which express a monoclonal antibody directed against oestriol.

[0175] Figure 30 is a plot showing positive results from an ELISA screen of phage displaying scFv fragments derived from the cell line 014 which express a monoclonal antibody directed against oestriol.

10 **[0176]** Figure 31 Electrophoresis of samples from stages of a Fab assembly. Samples from different stages in the PCR Fab assembly process described in example 26 were subjected to electrophoresis on a 1% TAE-agarose gel. Samples from a comparable scFv assembly process (as in example 14) are shown for comparison. Samples left to right are:

M = Markers

15 VHCH1 = sequences encoding VHCH1 domains amplified by PCR

VKCK = sequences encoding VKCK domains amplified by PCR

-L = Fab assembly reaction performed in absence of linker

+L = Fab PCR assembly reaction product VHCH1 plus VKCK plus linker

M = Markers

20 VK = sequences encoding VK domain amplified by PCR

VL = sequences encoding VH domains amplified by PCR

-L = scFv assembly reaction in absence of linker

+L = scFv assembly reaction in presence of linker

M = Markers

25

[0177] Figure 32. Comparison of ELISA signals with scFv D1.3 cloned in fd-CAT2 (fd) or pCAT-3. pCAT-3 scFv1.3 has been rescued with M13K07 (K07), M13K07ΔgIII No 3 (gIII No 3) or M13K07 gIIIΔNo 2 (g111No2). Phage antibodies are compared at 10 times (10x) 1 times (1x) or 0.1 times (0.1x) concentrations relative to concentration in the supernatant after overnight growth. The fdCAT2 and pCAT-3 non-recombinant vector signals were <0.01 at 10x concentration.

30

[0178] Figure 33. Western blot of PEG precipitated phage used in ELISA probed with anti-g3p. Free g3p and the g3p-scEvD1.3 fusion bands are arrowed.

Sample 1 - fd scFvD1.3

35 Sample 2 - pCAT3 vector

Sample 3 - pCAT3 scFvD1.3 rescued with M13K07, no IPTG

Sample 4 - pCAT3 scFvD1.3 rescued with M13K07, 50μM IPTG

Sample 5 - pCAT3 scFvD1.3 rescued with M13K07, 100μM IPTG

Sample 6 - pCAT3 scFvD1.3 rescued with M13K07 gIIIΔNo3 (no IPTG)

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Sample 7 - pCAT3 scFvD1.3 rescued with M13K07 gIIIΔ No 2 (no IPTG)

[0179] Panel A samples contain the equivalent of 8ul of phagemid culture supernatant per track, and 80μl of the fd supernatant (10-fold lower, phage yield than the phagemid). Panel B phagemid samples are those used in panel A at a five-fold higher sample loading (equivalent to 40μl of culture supernatant per track) to enable visualisation of the fusion band in samples rescued with parental M13K07.

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[0180] Figure 34 is a graph showing fdCAT2scFvD1.3 enrichment produced from a mixture of fdCAT2scFvD1.3 and fdCAT2TPB4 by one round of panning.

[0181] Figure 35 is a graph showing fdCAT2scFvD1.3 enrichment produced from a mixture of fdCAT2scFvD1.3 and fdCAT2TPB1 by one round of panning.

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[0182] Figure 36 shows the DNA sequence of scFv B18 (anti-NP).

[0183] Figure 37. shows a schematic representation of steps involved in the PCR assembly of nucleotide sequences encoding human Fab fragments. Details are in example 30.

[0184] Figure 38. shows A. a map of plasmid pJM1-FabD1.3 which is used for the expression of soluble human Fab fragments and as a template for the synthesis of linker DNA for Fab assembly. B. a schematic representation of sequences encoding a Fab construct. C. The sequence of DNA template for the synthesis of linker DNA for Fab assembly.

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[0185] Figure 39. shows a schematic representation of steps involved in the PCR assembly of nucleotide sequences encoding human scFv fragments. Details are in example 32.

[0186] Figure 40. ELISA assay of phage antibodies using plates coated with turkey egg lysogyme. Two clones B1

and A4 are shown derived by mutagenesis and selection from pAbD1.3 (example 35). Concentration (x axis) refers to the concentration of phage for each sample relative to the concentration in culture supernatant. B1 has raised binding to turkey egg lysogyme compared to D1.3. A4 has reduced binding to hen egg lysogyme compared to D1.3.

[0187] Figure 41. ELISA of phage antibodies binding to HEL and TEL. Clone 1 is fdCAT2scFvD1.3. Clones 2 to 10 were obtained from the library (example 36) after selection. The background values as defined by binding of these clones to BSA were subtracted.

[0188] Figure 42. shows the DNA sequence of the light chains D1.3 M1F and M21 derived by selection from a hierarchical library in example 36.

[0189] Figure 43 shows a Fv lambda expression vector (example 38) derived from pUC119. It contains the rearranged lambda germ line gene. The heavy and light chain cassettes each contain a ribosome binding site upstream of the pel B leader (Restriction sites shown as: H=Hind III; Sp=SphI; B=BamHI, E=EcoRI).

Materials and Methods

[0190] The following procedures used by the present applicants are described in Sambrook, J. et al., 1989 supra.: restriction digestion, ligation, preparation of competent cells (Hanahan method), transformation, analysis of restriction enzyme digestion products on agarose gels, purification of DNA using phenol/chloroform, 5'-end labelling of oligonucleotides, filter screening of bacterial colonies, preparation of 2xTY medium and plates, preparation of tetracycline stock solutions, PAGE of proteins, preparation of phosphate buffered saline.

[0191] All enzymes were supplied by New England Biolabs (CP Laboratories, PO Box 22, Bishop's Stortford, Herts., England) and were used according to manufacturer's instructions unless otherwise stated.

[0192] The vector fd-tet (Zacher, A.N. et al., 1980, supra) was obtained from the American Type Culture Collection (ATCC No. 37000) and transformed into competent TG1 cells (genotype: K12δ (lac-pro), sup E, thi, hsdD5/F traD36, pro A+B+, Lac 1^q, lac δM15).

[0193] Viral particles were prepared by growing TG1 cells containing the desired construct in 10 to 100 mls 2xTY medium with 15 µg/ml tetracycline for 16-24 hours. The culture supernatant was collected by centrifugation for 10 mins at 10,000 rpm in an 8 x 50 ml rotor, Sorval RC-5B centrifuge. Phage particles were precipitated by adding 1/5th volume 20% polyethylene glycol (PEG)/2.5M NaCl and leaving at 4°C for 1 hour. These were spun for 15 minutes as described above and the pellets resuspended in 10 mM Tris/HCl pH 8, 1mM EDTA to 1/100th of the original volume. Residual bacteria and undissolved material were removed by spinning for 2 minutes in a microcentrifuge. Single stranded DNA for mutagenesis or sequencing was prepared from concentrated phage according to Sambrook, J., et al., 1989,, supra.

Index of Examples

Example 1 Design of Insertion Point Linkers and Construction of Vectors

[0194] This example covers the construction of two derivatives of the phage vector fd-tet: a) fdTPs/Bs for the insertion of VH coding sequences; and b) fdTPs/Xh for the insertion of scFv coding sequences. The derivative vectors have a new BstEII site for insertion of sequences.

Example 2 Insertion of Immunoglobulin Fv Domain into Phage

[0195] This example covers the insertion of scFv coding sequences derived from an anti-lysozyme antibody D1.3 into fdTPs/Xh to give the construct fdTscFvD1.3.

Example 3 Insertion of Immunoglobulin VH Domain into Phage

[0196] This example covers the insertion of VH coding sequences derived from an anti-lysozyme antibody D1.3 into fdTPs/Bs to give the construct fdTVHD1.3.

Example 4 Analysis of Binding Specificity of Phage Antibodies

[0197] This example investigates the binding specificities of the constructs fdTscFvD1.3 and fdTVHD1.3.

Example 5 Construction of fdCAT2

[0198] This example covers the construction of the derivative fdCAT2 of the phage vector fdTPs/Xh. The derivative has restriction sites for enzymes that cut DNA infrequently.

Example 6 Specific Binding of Phage Antibody (pAb) to Antigen

[0199] This example shows the binding of pAb fdTscFvD1.3 to lysozyme by ELISA.

5 Example 7 Expression of FabD1.3

[0200] This example concerns the display of an antibody Fab fragment at the phage surface. The VH-CH1 chain is expressed by fdCAT2. The VL-CL chain is expressed by pUC19 in a bacterial host cell also infected with fdCAT2.

10 Example 8 Isolation of Specific, Desired Phage from a Mixture of Vector Phage

[0201] This example shows how a phage (e.g. fdTscFvD1.3) displaying a binding molecule can be isolated from vector phage by affinity techniques.

15 Example 9 Construction of pAb Expressing Anti-Hapten Activity

[0202] This example concerns the insertion of scFv coding sequences derived from the anti-oxazolone antibody NQ11 into fdTPs/Xh to generate the construct pAbNQ11. The example shows the binding of pAbNQ11 to oxazolone by ELISA.

20 Example 10 Enrichment of pAbD1.3 from Mixtures of other pAbs by Affinity Purification

[0203] This example shows how a phage (eg. pAbD1.3) displaying one sort of binding molecule can be isolated from phage (e.g. pAbNQ11) displaying another sort of binding molecule by affinity techniques.

25 Example 11 Insertion of Binding Molecules into Alternative Sites in the Phage

[0204] This example covers the insertion of scFv coding sequences derived from a) the anti-lysozyme antibody D1.3; and b) the anti-oxazolone antibody NQ11 into a BamHI site of fdTPs/Xh to give the constructs fdTBam1 having an NQ11 insert.

30 Example 12 PCR Assembly of Mouse VH and VLK Repertoires for Phage Display

[0205] This example concerns a system for the display on phage of all VH and VLK repertoires encoded by a mouse. The system involves the following steps. 1) Preparation of RNA from spleen. 2) Preparation of cDNA from the RNA 3) Use of primers specific for antibody sequences to PCR amplify all VH and VLK cDNA coding sequences 4) Use of PCR to create a linker molecule from linking pairs of VH and VLK sequences 5) Use of PCR to assemble continuous DNA molecules each comprising a VH sequence, a linker and a VLK sequence. The specific VH/VLK combination is randomly derived 6) Use of PCR to introduce restriction sites.

40 Example 13. Construction of Phagemid Containing Gene III Fused with the Coding Sequence for a Binding Molecule.
This example concerns the construction of two phagemids based on pUC119 which separately contain gene III from fdCAT2 and the gene III scFv fusion fdCAT2seFvD1.3 to generate pCAT2 and pCAT3 scFvD1.3 respectively.

45 Example 14. Rescue of Anti-Lysozyme Antibody Specificity from pCAT3scFvD1.3 by M13KO7

[0206] This example describes the rescue of the coding sequence for the gene IIIscFv fusion from pCAT3scFvD1.3 by M13M07 helper phage growth, phage were shown to be displaying scFv anti-lysozyme activity by ELISA.

50 Example 15. Transformation Efficiency of PCAT-3 and pCAT-3 scFvD1.3 Phagemids

[0207] This example compared the efficiency of the phagemids pVC119, pCAT-3 and pCAT3scFvD1.3 and the phage fdCAT2 to transform E.coli.

55 Example 16 PCR Assembly of a Single Chain Fv Library from an Immunised Mouse

[0208] This example concerns a system for the display on phage of scFv (comprising VH and VL) from. an immunised mouse using the basic technique outlined in example 14 (cDNA preparation and PCR assembly of the mouse VH and

VLK repertoires) and ligating the PCR assembled sequences into fdCAT2 to create a phage library of 10^5 clones. Testing of 500 clones showed that none showed specificity against phox.

Example 17 Selection of Antibodies Specific for 2-phenyl-5-oxazolone from a Repertoire from an Immunised Mouse.

[0209] This example shows that phage grown from the library established in example 16 can be subjected to affinity selection using phOx to select those phage displaying scFv with the desired specificity.

Example 18. Generation of Further Antibody Specificities by the Assembly of Hierarchical Libraries.

[0210] This example concerns the construction of hierarchical libraries in which a given VH sequence is combined with the complete VLK repertoire and a given VLK- sequence is combined with the complete VH repertoire and selection from these libraries of novel VH and VL pairings.

Example 19 Selection of Antibodies Displayed on Bacteriophage with Different Affinities for 2-phenyl-5-oxazolone using Affinity Chromatography

[0211] This example concerns the separation by affinity techniques of phages displaying scFv fragments with differing binding affinities for a given antigen.

Example 20. Construction of Phagemid pHEN1 for the Expression of Antibody Fragments Expressed on the surface of Bacteriophage following Superinfection

[0212] This example concerns the construction of the phagemid pHEN1 derived from pUC119. pHEN1 has the features shown in Fig. 26.

Example 21. Display of Single Chain Fv and Fab Fragments Derived from the Anti-Oxazolone Antibody NQ 10.12.5 on Bacteriophage fd using pHEN1 and fdCAT2.

[0213] This example describes the display of scFv and Fab fragment with a specificity against phOx on the surface of a bacteriophage. For display of scFv the phagemid pHEN1 comprises the sequences encoding scFv (VH and VL) for rescue by either the phages VSM13 or fdCAT2. For display of Fab the phage fdCAT2 comprises the sequence for either the H or L chain as a fusion with g3p and the phagemid pHEN1 comprises the sequence for the appropriate H or L chain partner.

Example 22. Rescue of Phagemid Encoding a Gene III protein Fusion with Antibody Heavy or Light Chains by Phage Encoding the Complementary Antibody Displayed on Phage and the Use of this Technique to make Dual Combinatorial Libraries

[0214] This example covers the use of phage antibodies encoding the antibody heavy or light chain to rescue a phagemid encoding a gene 3 protein fusion with the complementary chain and the assay of Fab fragments displayed on phage in ELISA. The use of this technique in the preparation of a dual combinatorial library is discussed.

Example 23. Induction of Soluble scFv and Fab Fragments using Phagemid pHEN1

[0215] This example covers the generation of soluble scFv and Fab fragments from gene III fusions with sequences encoding these fragments by expression of clones in pHEN1 in an E.coli strain which does not suppress amber mutations.

Example 24 Increased Sensitivity in ELISA of Lysozyme using fdTscFvD1.3 as Primary Antibody compared to Soluble scFvD1.3

[0216] This example covers the use of fdTscFvD1.3 in ELISA showing that lower amounts of lysozyme can be detected with phage antibody fdTscFvD1.3 than with soluble scFvD1.3.

Example 25 Direct Rescue and Expression of Mouse Monoclonal Antibodies as Single Chain Fv Fragments on the Surface of Bacteriophage fd

5 [0217] This example covers the display on phage as functional scFv fragments of two clones directly derived from cells expressing monoclonal antibodies directed against oestriol. Both clones were established to be functional using ELISA.

Example 26 PCR Assembly of DNA Encoding the Fab Fragment of an Antibody Directed against Oxazolone

10 [0218] This example covers the construction of a DNA insert encoding a Fab fragment by separate amplification of heavy and light chain DNA sequences followed by assembly. The construct was then inserted into the phage vector fdCAT2 and the phagemid vector pHEN1 and the Fab fragment displayed on the surface was shown to be functional.

Example 27 Construction of a Gene III Deficient Helper Phage

15 [0219] This example describes the construction of a helper phage derived from M13K07 by deleting sequences in gene III. Rescue of pCAT3-scFvD1.3 is described. The scFvD1.3 is expressed at a high level as a fusion using the deletion phage, equivalent to expression using fdCAT2-scFvD1.3.

20 Example 28 Selection of bacteriophage expressing scFv fragments directed against lysozyme from mixtures according to affinity using a panning procedure

25 [0220] This example concerns the selection of bacteriophage according to the affinity of the scFv fragment directed against lysozyme which is expressed on their surface. The phage of different affinities were bound to Petri dishes coated with lysozyme and, following washing, bound phage eluted using triethylamine. Conditions were found where substantial enrichment could be obtained for a phage with a 5-fold higher affinity than the phage with which it was mixed.

Example 29 Generation and Selection of Mutants of an Anti-4-hydroxy-3-nitrophenylacetic acid (NP) Antibody expressed on Phage using Mutator strains

30 [0221] This example covers the introduction of mutations into a gene for an antibody cloned in phage by growth of the phage in strains which randomly mutate DNA due to defects in DNA replication. Several mutations are introduced into phage which can then be selected from parent phage.

35 Example 30 A PCR Based Technique for one step Cloning of Human V-genes as Fab Constructs

[0222] This example gives methods for the assembly of Fab fragments from genes for antibodies. Examples are given for genes for antibodies directed against Rhesus-D in a human hybridoma and a polyclonal lymphoblastic cell line.

40 Example 31 Selection of Phage Displaying a Human Fab Fragment directed against the Rhesus-D Antigen by binding to Cells displaying the Rhesus D Antigen on their Surface

45 [0223] This example concerns the construction of, and display of phage antibodies from, a phagemid encoding a human Fab fragment directed against the Rhesus D antigen. Phage displaying this antigen were then affinity selected from a background of phage displaying scFvD1.3 anti-lysozyme on the basis of binding to Rhesus-D positive red blood cells.

Example 32 A PCR Based Technique for One Step Cloning of Human scFv Constructs

50 [0224] This example describes the generation of libraries of scFv fragments derived from an unimmunized human. Examples are given of the preparation for phage display of libraries in phagemids of scFv fragments derived from IgG and IgM sequences.

Example 33 Isolation of Binding Activities from a Library of scFvs from an Unimmunized Human

55 [0225] This example describes the isolation, from the library of scFv fragments derived from IgM genes of an unimmunized human, of clones for phage antibodies directed against BSA, lysozyme and oxazolone. Selection was by panning or affinity chromatography and analysis of binding specificity by ELISA. Sequencing of the clones showed

them to be of human origin.

Example 34 Rescue of human IgM library using helper phage lacking gene 3 ($\Delta g3$)

5 **[0226]** This example covers the isolation, from the library of scFv fragments of unimmunized human IgM genes, of clones of phage antibodies of clones for phage antibodies specific for thyroglobulin and oxazolone. In this example rescue was with M13K07gIII No3 (NCTC12478), a helper phage defective in gene III. Fewer rounds of selection appeared necessary for a phagemid library rescued with this phage compared to one rescued with M13K07.

10 Example 35 Alteration of Fine Specificity of scFvD1.3 displayed on Phage by Mutagenesis and Selection on Immobilized Turkey Lysozyme

[0227] This example covers the in vitro mutagenesis of pCATscFvD1.3 by replacement, with random amino acids, of residues known to be of importance in the preferential recognition of hen egg lysozyme over turkey egg lysozyme by scFvD1.3. Following selection for phage antibodies recognising turkey egg lysozyme by affinity chromatography, clones were analysed for specificity by ELISA. Two groups of clones were found with more equal recognition of hen and turkey lysozymes, one with increased ELISA signal with the turkey enzyme and one with reduced signal for the hen enzyme.

20 Example 36 Modification of the Specificity of an Antibody by Replacement of the VLK Domain by a VLK Library derived from an Unimmunised Mouse

[0228] This example shows that replacement of the VL domain of scFvD1.3 specific for hen eggwhite lysozyme (HEL) with a library of VL domains allows selection of scFv fragments which bind also to turkey eggwhite lysozyme (TEL). The scFv fragments were displayed on phage and selection by panning on tubes coated with TEL. Analysis by ELISA showed clones with enhanced binding to TEL compared to HEL. Those with highest binding to TEL were sequenced.

30 Example 37 Selection of a Phage Antibody Specificity by binding to an Antigen attached to Magnetic Beads. Use of a Cleavable Reagent to allow Elution of Bound Phage under Mild Conditions

[0229] This examples covers the use of a cleavable bond in the affinity selection method to allow release of bound phage under mild conditions. pAbNQ11 was enriched approximately 600 fold from a mixture with pAbD1.3 by selection using biotinylated Ox-BSA bound to magnetic beads. The cleavage of a bond between BSA and the biotin allows elution of the phage.

35 Example 38 Use of Cell Selection to provide an Enriched Pool of Antigen Specific Antibody Genes, Application to reducing the Complexity of Repertoires of Antibody Fragments Displayed on the Surface of Bacteriophage

40 **[0230]** This example covers the use of cell selection to produce an enriched pool of genes encoding antibodies directed against 4-hydroxy-3-nitrophenylacetic acid and describes how this technique could be used to reduce the complexity of antibody repertoires displayed on the surface of bacteriophage.

Example 1

45 Design of Insertion Point Linkers and Construction of Vectors

[0231] The vector fd-tet has two BstEII restriction sites flanking the tetracycline resistance gene (fig 3). Since the strategy for inserting the VH fragments was to ligate them into a newly inserted BstEII site within gene III, it was advantageous to delete the original BstEII sites from fd-tet. This was achieved by digesting fd-tet with the restriction enzyme BstEII, filling-in the 5' overhangs and re-ligating to generate the vector fdT δ Bst. Digestion of fd-tet with BstEII (0.5 units/ μ l) was carried out in 1x KGB buffer (100 mM potassium glutamate, 23 mM Tris-acetate (pH 7.5), 10 mM magnesium acetate, 50 μ g/ml bovine serum albumin, 0.5 mM dithiothreitol (Sambrook, J., et al., 1989, supra.) with DNA at a concentration of 25 ng/ μ l. The 5' overhang was filled in, using 2x KGB buffer, 250 μ M each dNTP's (Pharmacia Ltd., Pharmacia House, Midsummer Boulevard, Milton Keynes, Bucks., UK.) and Klenow Fragment (Amersham International, Lincoln Place, Green End, Aylesbury, Bucks., UK) at 0.04 units/ μ l. After incubating for 1 hour at room temperature, DNA was extracted with phenol/chloroform and precipitated with ethanol.

[0232] Ligations were carried out at a DNA concentration of 50ng/ μ l. Ligations were transformed into competent TG1 cells and plated onto TY plates supplemented with 15 μ g/ml tetracycline. This selects for vectors where the gene

for tetracycline resistance protein has reinserted into the vector during the ligation step. Colonies were picked into 25 mls of 2xTY medium supplemented with 15 µg/ml tetracycline and grown overnight at 37°C.

[0233] Double stranded DNA was purified from the resulting clones using the gene-clean II kit (Bio101 Inc., PO Box 2284, La Jolla, California, 92038-2284, USA.) and according to the small scale rapid plasmid DNA isolation procedure described therein. The orientation of 5 of the resulting clones was checked using the restriction enzyme *Cla*I. A clone was chosen which gave the same pattern of restriction by *Cla*I as fd-tet, but which had no *Bst*E II sites.

[0234] In vitro mutagenesis of fdTδBst was used to generate vectors having appropriate restriction sites that facilitate cloning of antibody fragments downstream of the gene III signal peptide and in frame with the gene III coding sequence. The oligonucleotide directed mutagenesis system version 2 (Amersham International) was used with oligo 1 (figure 4) to create fdTPs/Bs (to facilitate cloning of VH fragments). The sequence of fdTPs/Bs (figure 4) was confirmed using the sequenase version 2.0 kit (USB Corp., PO Box 22400, Cleveland, Ohio, 44122, USA.) with oligo 3 (figure 4) as a primer.

[0235] A second vector fdTPs/Xh (to facilitate cloning of single chain Fv fragments) was generated by mutagenising fdTPs/Bs with oligo 2 according to the method of Venkitaraman, A.R., Nucl. Acid Res. 17, p 3314. The sequence of fdTPs/Xh (figure 4) was confirmed using the sequenase version 2.0 kit (USB Corp.) with oligo 3 as a primer.

[0236] Clearly, alternative constructions will be apparent to those skilled in the art. For example, M13 and/or its host bacteria could be modified such that its gene III could be disrupted without the onset of excessive cell death.

[0237] Moreover, the invention employs a plasmid containing a single stranded phage replication origin, such as pUC119, so that where modified fd gene III, or other modified protein, is incorporated into the plasmid, superinfection with modified phage such as K07 results in the encapsulation of the phage antibody genome in a coat partially derived from the helper phage and partly from the phage antibody gene III construct.

[0238] The detailed construction of a vector such as fdTPs/Bs is only one way of achieving the end of a phage antibody. For example, techniques such as sticky feet cloning/mutagenesis (Clackson, T. and Winter, G. 1989 Nucl. Acids. Res., 17, p 10163-10170) could be used to avoid use of restriction enzyme digests and/or ligation steps.

Example 2.

Insertion of Immunoglobulin Fv Domain into Phage

[0239] The plasmid scFv D1.3 myc (gift from G. Winter and A. Griffiths) contains VH and VL sequences from the antibody D1.3 fused via a peptide linker sequence to form a single chain Fv version of antibody D1.3. The sequence of the scFv and surrounding sequences in scFvD1.3 myc is shown in figure 5.

[0240] The D1.3 antibody is directed against hen egg lysozyme (Harper, M. et al., 1987, Molec. Immunol. 24, 97-108) and the scFv form expressed in E.coli has the same specificity (A. Griffiths and G. Winter personal Communication).

[0241] Digestion of scFv D1.3 myc with *Pst*I and *Xho*I (these restriction sites are shown on Fig. 5), excises a fragment of 693 bp which encodes the bulk of the scFv. Ligation of this fragment into fdTPs/Xh cleaved with *Pst*I and *Xho*I gave rise to the construct fdTscFvD1.3 encoding the gene III signal peptide and first amino acid fused to the complete D1.3 scFv, followed by the mature gene III protein from amino acid 2.

[0242] The vector fdTPs/Xh was prepared for ligation by digesting with the *Pst*I and *Xho*I for 2 hours followed by digestion with calf intestinal alkaline phosphatase (Boehringer Mannheim UK Ltd., Bell Lane, Lewes, East Sussex, BN7 1LG) at one unit/ul for 30 minutes at 37°C. Fresh calf intestinal alkaline phosphatase was added to a final total concentration of 2 units/ul and incubated for a further 30 minutes at 37°C. The reaction was extracted three times with phenol/chloroform, precipitated with ethanol and dissolved in water. The insert from scFvD1.3 myc was excised with the appropriate restriction enzymes (*Pst*I and *Xho*I) extracted twice with phenol/chloroform, precipitated with ethanol and dissolved in water. Ligations were carried out as described in example 1, except both vector and insert samples were at a final concentration of 5 ng/ul each. The formation of the correct construct was confirmed by sequencing as described in example 1.

[0243] To demonstrate that proteins of the expected size were produced, virions were concentrated by PEG precipitation as described above. The samples were prepared for electrophoresis as described in Sambrook J. et al 1989 supra. The equivalent of 2mls of supernatant was loaded onto an 18% SDS polyacrylamide gel. After electrophoresis, the gel was soaked in gel running buffer (50 mM Tris, 380 mM Glycine, 0.1% SDS) with 20% methanol for 15 minutes. Transfer to nitrocellulose filter was executed in fresh 1x running buffer/20% methanol using TE70 Semi Phor a semi-dry blotting apparatus (Hoeffer, 654 Minnesota Street, Box 77387, San Francisco, California 94107, USA.).

[0244] After transfer, the filter was blocked by incubation for 1 hour in a 2% solution of milk powder (Marvel) in phosphate buffered saline (PBS). Detection of scFv and VH protein sequences in the phage antibody fusion proteins was effected by soaking the filter for 1 hour with a 1/1000 dilution (in 2% milk powder) of a rabbit polyclonal antiserum raised against affinity purified, bacterially expressed scFv fragment (gift from G. Winter). After washing with PBS (3 x 5 minute washes), bound primary antibody was detected using an anti-rabbit antibody conjugated to horseradish per-

oxidase (Sigma, Fancy Road, Poole, Dorset, BH17 7NH, UK.) for 1 hour. The filter was washed in PBS/0.1% triton X-100 and developed with 0.5 mg/ml 3,3'-diaminobenzidine tetrahydrochloride (DAB), 0.02% cobalt chloride, 0.03% hydrogen peroxide in PBS.

[0245] The results showed that with clones fdTVHD1.3 (from example 3 incorporating sequences coding for VH) and fdTscFvD1.3 (incorporating sequences coding for scFv) a protein of between 69,000 and 92,500 daltons is detected by the anti-Fv serum. This is the expected size for the fusion proteins constructed. This product is not observed in supernatants derived from fd-tet, fdTδBst or fdTPs/Xh.

Example 3.

Insertion of Immunoglobulin VH Domain into Phage Antibody

[0246] The VH fragment from D1.3 was generated from the plasmid pSW1-VHD1.3-TAG1 (Ward, E.S. et al., 1989 supra.). Digestion of this plasmid with Pst1 and BstEII generates the fragment shown between positions 113 and 432 in figure 5. Cloning of this fragment into the Pst1 and BstEII sites of fdTPs/Bs gave rise to the construct fdTVHD1.3 which encodes a fusion protein with a complete VH domain inserted between the first and third amino acids of the mature gene III protein (amino acid two has been deleted).

[0247] The methods used were exactly as in example 2 except that the vector used was fdTPs/Bs digested with Pst1 and BstEII.

Example 4.

Analysis of Binding Specificity of Phage Antibodies

[0248] The binding of the various phage antibodies to the specific antigen, lysozyme, was analysed using ELISA techniques. Phage antibodies (e.g. fdTVHD1.3 and fdTscFvD1.3) were grown in E.coli and Phage antibody particles were precipitated with PEG as described in the materials and methods. Bound phage antibody particles were detected using polyclonal sheep serum raised against the closely related phage M13.

[0249] ELISA plates were prepared by coating 95 well plates (Falcon Microtest III flexible plate. Falcon: Becton Dickinson Labware, 1950 Williams Drive, Oxnard, California, 93030, USA.) with 200 ul of a solution of lysozyme (1mg/ml unless otherwise stated) in 50 mM NaHCO₃ for 16-24 hours. Before use, this solution was removed, the plate rinsed several times in PBS and incubated with 200 ul of 2% milk powder/PBS for 1 hour. After rinsing several times with PBS, 100 ul of the test samples were added and incubated for 1 hour. Plates were washed (3 rinses in 0.05% Tween 20/PBS followed by 3 rinses in PBS alone). Bound phage antibodies were detected by adding 200 ul/well of a 1/1000 dilution of sheep anti-M13 polyclonal antiserum (gift from G. Winter, although an equivalent antibody can be readily made by one skilled in the art using standard methodologies) in 2% milk powder/PBS and incubating for 1 hour. After washing as above, plates were incubated with biotinylated anti-sheep antibody (Amersham International) for 30 minutes. Plates were washed as above, and incubated with streptavidinhorseradish peroxidase complex (Amersham International). After a final wash as above, 0.5 mg/ml ABTS substrate in citrate buffer was added (ABTS = 2'2'-azinobis (3-ethylbenzthiazoline sulphonic acid); citrate buffer = 50 mM citric acid, 50 mM tri-sodium citrate at a ratio of 54:46. Hydrogen peroxide was added to a final concentration of 0.003% and the plates incubated for 1 hour. The optical density at 405 nm was read in a Titertek multiskan plate reader.

[0250] Figure 6 shows the effect of varying the amount of phage antibody. 100 ul of various dilutions of PEG precipitated phage were applied and the amount expressed in terms of the original culture volume from which it was derived. Signals derived from both the scFv containing phage antibody (fdTscFvD1.3) and the VH containing phage antibody (fdTVHD1.3) and the VH containing phage antibody were higher than that derived from the phage antibody vector (fdTPs/Xh). The highest signal to noise ratio occurs using the equivalent of 1.3 mls of culture.

[0251] Figure 7 shows the results of coating the plates with varying concentrations of lysozyme or bovine serum albumin (BSA). The equivalent of 1 ml of the original phage antibody culture supernatant was used. The signals from supernatants derived from fdTscFvD1.3 were again higher than those derived from fdTPs/Xh when lysozyme coated wells were used. There was no significant difference between these two types of supernatant when the plates were coated with BSA. Broadly speaking the level of signal on the plates is proportional to the amount of lysozyme coated. These results demonstrate that the binding detected is specific for lysozyme as the antigen.

Example 5.Construction of fd CAT 2

[0252] It would be useful to design vectors that enable the use of restriction enzymes that cut DNA infrequently, thus avoiding unwanted digestion of the antibody gene inserts within their coding sequence. Enzymes with an eight base recognition sequence are particularly useful in this respect, for example Not1 and Sfil. Chaudhary et al (PNAS 87 p1066-1070, 1990) have identified a number of restriction sites which occur rarely in antibody variable genes. The applicant has designed and constructed a vector that utilises two of these sites, as an example of how this type of enzyme can be used. Essentially sites for the enzymes ApaL1 and Not1 were engineered into fdTPs/Xh to create fdCAT2.

The oligonucleotide:

5' ACT TTC AAC AGT TTC TGC GGC CGC CCG TTT GAT CTC GAG CTC
CTG CAG TTG GAC CTG TGC ACT GTG AGA ATA GAA 3'

was synthesised (supra fig 4 legend) and used to mutagenise fdTPs/Xh using an in vitro mutagenesis kit from Amersham International as described in example 1, to create fd-CAT2. The sequence of fd-CAT2 was checked around the site of manipulation by DNA sequencing. The final sequence around the insertion point within gene III is shown in figure 8.

N.B. fdCAT2 is also referred to herein by the alternative terminologies fd-tet-DOG1 and fdDOG1.

Example 6Specific Binding of Phage-antibody (pAb) to Antigen

[0253] The binding of pAb D1.3 (fdTscFvD1.3 of example 2) to lysozyme was further analysed by ELISA.

Methods.1. Phage growth.

[0254] Cultures of phage transduced bacteria were prepared in 10-100 mls 2 x TY medium with 15 µg/ml tetracycline and grown with shaking at 37°C for 16-24 hrs. Phage supernatant was prepared by centrifugation of the culture (10 min at 10,000 rpm, 8 x 50 ml rotor, Sorval RC-5B centrifuge). At this stage, the phage titre was $1 - 5 \times 10_{10}$ /ml transducing units. The phage were precipitated by adding 1/5 volume 20% PEG 2.5 M NaCl, leaving for 1 hr at 4°C, and centrifuging (supra). The phage pellets were resuspended in 10 mM Tris-HCl, 1mM EDTA pH 8.0 to 1/100th of the original volume, and residual bacteria and aggregated phage removed by centrifugation for 2 min in a bench microcentrifuge.

ELISA

[0255] Plates were coated with antigen (1 mg/ml antigen) and blocked as described in example 4. $2 \times 10_{10}$ phage transducing units were added to the antigen coated plates in phosphate buffered saline (PBS) containing 2% skimmed milk powder (MPBS). Plates were washed between each step with three rinses of 0.5% Tween-20 in PBS followed by three rinses of PBS. Bound phage was developed by incubating with sheep anti-M13 antisera and detected with horseradish peroxidase (HRP) conjugated anti-goat serum (Sigma, Poole, Dorset, UK) which also detects sheep immunoglobulins and ABTS (2'2'-azinobis (3-ethylbenzthiazoline sulphonic acid). Readings were taken at 405 nm after a suitable period. The results (figure 9) show that the antibody bearing-phage had the same pattern of reactivity as the original D1.3 antibody (Harper, M., Lema, F., Boulot, G., and Poljak, F.J. (1987) Molec. Immunol. 24, 97-108), and bound to hen egg-white lysozyme, but not to turkey egg-white lysozyme, human lysozyme or bovine serum albumin. The specificity of the phage is particularly illustrated by the lack of binding to the turkey egg-white lysozyme that differs from hen egg-white lysozyme by only 7 amino acids.

Example 7.Expression of Fab D1.3

[0256] The aim of this example was to demonstrate that the scFv format used in example 2 was only one way of displaying antibody fragments in the pAb system. A more commonly used antibody fragment is the Fab fragment (figure 1) and this example describes the construction of a pAb that expresses a Fab-like fragment on its surface and shows that it binds specifically to its antigen. The applicant chose to express the heavy chain of the antibody fragment consisting of the VH and CH1 domains from coding sequences within the pAb itself and to coexpress the light chain in the bacterial host cell infected with the pAb. The VH and CH1 regions of anti-lysozyme antibody D1.3 were cloned in fd CAT2, and the corresponding light chain cloned in plasmid pUC19. The work of Skerra and Pluckthun (Science 240, p1038-1040 (1988) and Better et al 1988 supra; demonstrated that multimeric antigen binding fragments of the antibody molecule could be secreted into the periplasm of the bacterial cell in a functional form using suitable signal sequences. However, in these publications, special measures were described as being needed to recover the binding protein from the cell, for example Skerra and Pluckham needed to recover the Fv fragment from the periplasm by affinity chromatography. The present applicants have shown that it is possible to direct the binding molecule to the outside of the cell on a phage particle, a process that requires several events to occur: correct secretion and folding of the binding molecule; association of the chains of the binding molecule; correct assembly of the phage particle; and export of the intact phage particle from the cell.

[0257] Alternatively, it is possible however, to express the light chain from within the pAb genome by, for example, cloning an expression cassette into a suitable place in the phage genome. Such a suitable place would be the intergenic region which houses the multicloning sites engineered into derivative of the related phage M13 (see, for example, Yanisch-Perron, C. et al., Gene 33, p103-119, (1985)).

[0258] The starting point for this example was the clone Fab D1.3 in pUC19, a map of which is shown in figure 10. The regions hybridising with the oligonucleotides KSJ6 and 7 below are shown underlined in fig 10. The sequence encoding the VH-CH1 region (defined at the 5' and 3' edges by the oligonucleotides KSJ6 and 7 below) was PCR amplified from Fab D1.3 in pUC19 using oligonucleotides KSJ 6 and 7, which retain the Pst I site at the 5' end and introduce a Xho I site at the 3' end, to facilitate cloning into fd CAT2. The sequences for the oligonucleotides KSJ6 and 7 are shown below. The underlined region of KSJ7 shows the portion hybridising with the sequence for D1.3.

KSJ6: 5' AGG TGC AGC TGC AGG AGT CAG G 3'

KSJ7: 5' GGT GAC CTC GAG TGA AGA TTT GGG CTC AAC TTT C 3' PCR conditions were as described in example II, except that thirty cycles of PCR amplification were performed with denaturation at 92°C for 45 seconds, annealing at 55°C for 1 minute and extension at 72°C for 1 minute. The template used was DNA from TG1 cells containing Fab D1.3 in pUC19 resuspended in water and boiled. The template DNA was prepared from the colonies by picking some colony material into 100µl of distilled H₂O and boiling for 10 mins. 1µl of this mixture was used in a 20µl PCR. This regime resulted in amplification of the expected fragment of approximately 600bp. This fragment was cut with Pst I and Xho I, purified from an agarose gel and ligated into Pst I/Xho I-cut fdCAT2. The PCR mixture was extracted with phenol/chloroform and ethanol precipitated (Sambrook et al. supra.) before digestion with Pst I and Xho I (New England Biolabs according to manufacturers recommendations. The fragment was resolved on 1% Tris-Acetate EDTA agarose gel (Sambrook et al. supra) and purified using Geneclean (BIO 101, Geneclean, La Jolla, San Diego, California, USA) according to manufacturers recommendations.

[0259] fd-CAT2 vector DNA was digested with Pst I and Xho I (New England BioLabs) according to manufacturers recommendations, extracted with phenol/chloroform and ethanol precipitated (Sambrook et al. supra.).

[0260] 75ng of Pst I/Xho I-digested vector DNA was ligated to 40ng of PCR-amplified Pst I/Xho I-digested hEGF-R fragment in 12µl of ligation buffer (66mM TrisHCl (pH7.6), 5mM MgCl₂, 5mM dithiothreitol, (100µg/ml bovine serum albumin, 0.5mM ATP, 0.5mM Spermidine) and 400 units T4 DNA ligase (New England BioLabs) for 16 hours at 16°C.

[0261] Two µl of the ligation mixture was transformed into 200µl of competent E.coli MC1061 cells, plated on 2TY agar containing 15µg/ml tetracycline and incubated at 30°C for 20 hours. A portion of the ligation reaction mixture was transformed into E.coli MC1061 (Available from, for example Clontech Laboratories Inc, Palo Alto, California) and colonies identified by hybridisation with the oligonucleotide D1.3CDR3A as described in example 10. The presence of the VHCH1 gene fragment was likewise confirmed by PCR, using oligonucleotides KSJ6 and 7. A representative clone was called fd CAT2VHCH1 D1.3. The heavy-chain was deleted from Fab D1.3 in pUC19 by Sph I cleavage of Fab D1.3 plasmid DNA. The pUC 19 2.7Kb fragment containing the light chain gene was purified from a TAE agarose gel, and 10ng of this DNA self-ligated and transformed into competent E.coli TG1. Cells were plated on 2TY agar containing ampicillin (100µg/ml) and incubated at 30°C overnight. The resulting colonies were used to make miniprep DNA (Sambrook et al. supra), and the absence of the heavy chain gene confirmed by digestion with Sph I and Hind III. A representative clone was called LCD1.3 DHC.

[0262] An overnight culture of fd CAT2VHCH1 D1.3 cells was microcentrifuged at 13,000Xg for 10 minutes and 50µl

of the supernatant containing phage particles added to 50 µl of an overnight culture of LCD1.3 DHC cells. The cells were incubated at 37°C for 10 minutes and plated on 2TY agar containing ampicillin (100 µg/ml) and 15 µg/ml tetracycline. Phage were prepared from some of the resulting colonies and assayed for their ability to bind lysozyme as described in example 6.

[0263] The results (Figure 11) showed that when the heavy and light chain Fab derivatives from the original antibody D1.3 were present, the pAb bound to lysozyme. pAb expressing the fd VHCH1 fragment did not bind to lysozyme unless grown in cells also expressing the light chain. This shows that a functional Fab fragment was produced by an association of the free light chain with VHCH1 fragment fused to gene III and expressed on the surface of the pAb.

Example 8

Isolation of Specific, Desired Phage from a Mixture of Vector Phage.

[0264] The applicant purified pAb (D1.3) (originally called fdTscFvD1.3 in example 2) from mixtures using antigen affinity columns. pAb (D1.3) was mixed with vector fd phage (see table 1) and approximately 10^{12} phage passed over a column of lysozyme-Sepharose (prepared from cyanogen bromide activated sepharose 4B (Pharmacia, Milton Keynes, Bucks, UK.) according to the manufacturers instructions. TG1 cells were infected with appropriate dilutions of the elutes and the colonies derived, were analysed by probing with an oligonucleotide that detects only the pAb (D1.3) see Table 1 and Fig. 12. A thousand fold enrichment of pAb(D1.3) was seen with a single column pass. By growing the enriched phage and passing it down the column again, enrichments of up to a million fold were seen.

[0265] Enrichment was also demonstrated using purely immunological criteria. For example, 10^{12} phage (at a ratio of 1 pAb (D1.3) to 4×10^6 fdTPs/Bs) was subjected to two rounds of affinity selection, and then 26 colonies picked and grown overnight. The phage was then assayed for lysozyme binding by ELISA (as example 6). Five colonies yielded phage with lysozyme binding activities, see table 1, and these were shown to encode the scFv (D1.3) by PCR screening (example 13, using 30 cycles of 1 minute at 92°C, 1 minute at 60°C, 1 minute at 72°C using CDR3PCR1 and oligo 3 (fig. 4) as primers).

[0266] Thus very rare pAbs can be fished out of large populations, by using antigen to select and then screen the phage.

[0267] In this example, affinity chromatography of pAbs and oligonucleotide probing were carried out as described below.

[0268] Approximately 10^{12} phage particles in 1 ml MPBS were loaded onto a 1 ml lysozyme-Sepharose affinity column which had been prewashed in MPBS. The column was washed in turn with 10 ml PBS; then 10 ml 50 mM Tris-HCl, 500 mM NaCl pH 7.5; then 10 ml 50 mM Tris-HCl 500 mM NaCl pH 8.5; then 5 ml 50 mM Tris-HCl, 500 mM NaCl pH 9.5 (adjusted with triethylamine) and then eluted with 5 ml 100 mM triethylamine. The eluate was neutralised with 0.5 M sodium phosphate buffer pH 6.8 and the phage plated for analysis. For a second round of affinity chromatography, the first column eluate was plated to about 30,000 colonies per petri dish. After overnight growth, colonies were then scraped into 5 ml 2 x TY medium, and a 20 µl aliquot diluted into 10 ml fresh medium and grown overnight. The phage was PEG precipitated as described above, resuspended in 1 ml MPBS and loaded onto the column, washed and eluted as above.

Oligonucleotides synthesised:

CDR3PCR1 5'TGA GGA C(A or T) C(A or T) GC CGT CTA CTA CTG
TGC 3'

[0269] 40 pmole of oligonucleotide VH1FOR (Ward, E. S., et al (1989) Nature 341, 544-546), specific to pAb (D1.3) was phosphorylated with 100 µCi α -³²P ATP, hybridised (1 pmole/ml) to nitrocellulose filters at 67°C in 6 x saline sodium citrate (SSC) Sambrook et al., supra. buffer for 30 minutes and allowed to cool to room temperature for 30 mins, washed 3 x 1 min at 60°C in 0.1 x SSC.

Example 9

Construction of pAb Expressing Anti-hapten Activity

[0270] Oxazolone is a hapten that is commonly used for studying the details of the immune response. The anti-oxazolone antibody, NQ11 has been described previously (E. Gherardi, R. Pannell, C. Milstein, J. Immunol. Method 126 61-68). A plasmid containing the VH and VL gene of NQ11 was converted to a scFv form by inserting the BstEI/SacI fragment of scFvD1.3 myc (nucleotides 432-499 of Fig. 5) between the VH and VL genes to generate pscFvNQ11,

the sequence of which is shown in fig. 13. This scFv was cloned into the Pst1/Xho1 site of FdTPs/Xh (as described earlier) to generate pAb NQ11 has an internal Pst1 site and so it was necessary to do a complete digest of pscFvNQ11 with Xho1 followed by a partial digest with Pst1).

[0271] The specific binding of pAb NQ11 was confirmed using ELISA. ELISA plates were coated at 37°C in 50 mM NaHCO₃ at a protein concentration of 200 µg/ml. Plates were coated with either hen egg lysozyme (HEL), bovine serum albumin (BSA), or BSA conjugated to oxazolone (OX-BSA) (method of conjugation in Makela O., Kartinen M., Pelkonen J.L.T., Karjalainen K. (1978) J. Exp. Med. 148 1644). Preparation of phage, binding to ELISA plates, washing and detection was as described in example 6. Samples were assayed in duplicate and the average absorbance after 10 minutes presented in figure 14.

[0272] This result demonstrates that the pAb NQ11 binds the correct antigen. Figure 14 also shows that pAb D1.3 and pAb NQ11 bind only to the antigen against which the original antibodies were raised.

Example 10

Enrichment of pAb D1.3 from Mixtures of Other pAb by Affinity Purification

[0273] 3 x 10¹⁰ phage in 10 mls of PBSM at the ratios of pAb D1.3 to pAb NQ11 shown in table 2 were passed over a 1 ml lysozyme Sepharose column. Washing, elution and other methods were as described in example 8 unless otherwise stated. Eluates from the columns were used to infect TG1 cells which were then plated out. Colonies were probed with a probe which distinguishes pAb D1.3 from pAb NQ11. The sequence of this oligonucleotide (D1.3CDR3A) is:-

5'GTA GTC AAG CCT ATA ATC TCT CTC 3'

[0274] Table 2 presents the data from this experiment. An enrichment of almost 1000 fold was achieved in one round and an enrichment of over a million fold in two rounds of purification. This parallels the result described in example 8.

Example 11

Insertion of Binding Molecules into Alternative Sites in the Phage

[0275] The availability of an alternative site in the phage for the insertion of binding molecules would open up the possibility of more easily expressing more than one binding molecule e.g. an antibody fragment in a single pAb. This may be used to generate single or multiple binding specificities. The presence of two distinct binding activities on a single molecule will greatly increase the utility and specificity of this molecule. It may be useful in the binding of viruses with a high mutational rate such as human immunodeficiency virus. In addition, it may be used to bring antigens into close proximity (e.g. drug targeting or cell fusion) or it may act as a "molecular clamp" in chemical, immunological or enzymatic processes.

[0276] The vector fd-tet and the derivatives described here, have a single BamH1 site in gene 3. This has previously been used for the expression of peptide fragments on the surface of filamentous bacteriophage (Smith GP. (1985) Science 228 p1315-1317 and de la Cruz et al. (1988) J Biol. Chem. 263 p4318-4322). This provides a potential alternative site for the insertion of antibody fragments.

[0277] DNA fragments encoding scFv's from D1.3 or NQ11 were generated by PCR using the primers shown below. These primers were designed to generate a fragment with BamH1 sites near both the termini, to enable cloning into the BamH1 site of gene3 (see figure 16(1)). The oligonucleotides used, also ensure that the resulting PCR product lacks Pst1 and Xho1 restriction sites normally used for manipulating the scFv's (see figure 16(1)). This will facilitate subsequent manipulation of a second antibody fragment in the usual way at the N terminus of gene 3. The oligonucleotides used were:-

G3Bam1 5' TTT AAT GAG GAT CCA CAG GTG CAG CTG CAA GAG 3'
G3Bam2 5' AAC GAA TGG ATC CCG TTT GAT CTC AAG CTT 3'.

Preparation of vector and PCR insert

[0278] The PCR reaction was carried out in an 80 µl reaction as described in example 11 using 1ng/µl of template and 0.25U/µl of Taq polymerase and a cycle regime of 94°C for 1 minute, 60°C for 1 minute and 70°C for 2 minutes

over 30 cycles. The template was either pscFvNQ11 (example 9) or scFvD1.3 myc (example 2). Reaction products were extracted with phenol:chloroform, precipitated, dissolved in water and digested with BamH1 according to manufacturers instructions. The digest was re-extracted with phenol: chloroform, precipitated and dissolved in water.

[0279] The vector fdTPs/Xh was cleaved with BamH1 and treated with calf intestinal phosphatase and purified as described in example 2. Ligations were set up at a vector concentration of approximately 6ng/μl and a PCR insert concentration of approximately 3ng/μl; These were ligated for 2.5 hours at room temperature before transforming into competent TG1 cells and plating on TY tet plates. The resultant colonies were probed as described in example 8. DNA was prepared from a number of colonies and the correct orientation and insert size confirmed by restriction digestion with Hind III in isolation or in combination with BamH1. (One Hind III site is contributed by one of the primers and the other by the vector).

[0280] Two clones containing a D1.3 insert (fdTBaml) and fdTBam2) and one containing an NQ11 insert (NQ11Baml) were grown up and phage prepared as described earlier. ELISAs were carried out as described in example 6. No specific signal was found for any of these clones suggesting that the natural BamH1 site is not a suitable site for insertion of a functional antibody (results not shown).

[0281] It may be possible to clone, into alternative sites to retain binding activity. The peptide repeats present in gene III may provide such a site (figure 16 blocks A and B). This can be done by inserting a BamH1 site and using the PCR product described above. To facilitate this, the natural BamH1 site was removed by mutagenesis with the oligonucleotide G3mutδBam shown below (using an in vitro mutagenesis kit (Amersham International)):-

G3mutδBam 5' CA AAC GAA TGG GTC CTC CTC ATT A 3'

The underlined residue replaces an A residue, thereby removing the BamH1 site. DNA was prepared from a number of clones and several mutants lacking BamH1 sites identified by restriction digestion.

[0282] The oligonucleotide G3 Bamlink was designed to introduce a BamH1 site at a number of possible sites within the peptide linker sites A and B, see figure 16(2). The sequence of the linker is:
Bamlink 5'CC (G or A) CC ACC CTC GGA TCC (G or A) CC ACC CTC 3'
Its relationship to the peptide repeats in gene III is shown in figure 16.

Example 12

PCR Assembly of Mouse VH and VL Kappa (VLK) Repertoires for Phage Display

[0283] The principle is illustrated in figure 17. Details are provided in sections A to F below but the broad outline is first discussed.

1. cDNA is prepared from spleen RNA from an appropriate mouse and the VH and VLK repertoires individually amplified. Separately, primers reverse and complementary to VH1FOR-2 (domain 1) and VLK2BACK (domain 2) are used to amplify an existing scFv-containing DNA by PCR. (The term FOR refers to e.g. a primer for amplification of sequences on the sense strand resulting in antisense coding sequences. The term BACK refers to e.g. a primer for amplification of sequences on the antisense strand resulting in sense coding sequences). This generates a 'linker' molecule encoding the linker with the amino acid sequence (1 letter code) (GGGGS)₃ which overlaps the two primary (VH and VLK) PCR products.

2. The separate amplified VH, VLK and linker sequences now have to be assembled into a continuous DNA molecule by use of an 'assembly' PCR. In the secondary 'assembly' PCR, the VH, VLK and linker bands are combined and assembled by virtue of the above referred to overlaps. This generates an assembled DNA fragment that will direct the expression of VH and one VLK domain. The specific VH/VLK combination is derived randomly from the separate VH and VLK repertoires referred to above.

[0284] The assembly PCR is carried out in two stages. Firstly, 7 rounds of cycling with just the three bands present in the PCR, followed by a further 20 rounds in the presence of the flanking primers VH1BACK (referring to domain 1 of VH) and VLKFOR. The nucleotide sequences for these oligonucleotide primers are provided under the section entitled 'Primer Sequences' below. This two stage process, avoids the potential problem of preferential amplification of the first combinations to be assembled.

[0285] For cloning into the phage system, the assembled repertoires must be 'tagged' with the appropriate restriction sites. In the example provided below this is illustrated by providing an ApaI1 restriction site at the VH end of the continuous DNA molecule and a Not 1 site at the VLK end of the molecule. This is carried out by a third stage PCR using tagged primers. The nucleotide sequences for these oligonucleotide primers are also provided under the section

entitled 'Primer Sequences' below. There are however, 4 possible kappa light chain sequences (whereas a single consensus heavy chain sequence can be used). Therefore 4 oligonucleotide primer sequences are provided for VLK.

[0286] For this third stage PCR, sets of primers which create the new restriction site and have a further 10 nucleotides on the 5' side of the restriction site have been used. However, long tags may give better cutting, in which case 15-20 nucleotide overhangs could be used.

[0287] Scrupulously clean procedures must be used at all times to avoid contamination during PCR. Negative controls containing no DNA must always be included to monitor for contamination. Gel boxes must be depurinated. A dedicated GeneClean kit (B10 101, GeneClean, La Jolla, San Diego, California, USA) can be used according to manufacturers instructions to extract DNA from an agarose gel. The beads, NaI and the NEW wash should be aliquoted.

[0288] All enzymes were obtained from CP Laboratories, P.O. Box 22, Bishop's Stortford, Herts CM20 3DH and the manufacturers recommended and supplied buffers were used unless otherwise stated.

A. RNA Preparation

[0289] RNA can be prepared using many procedures well known to those skilled in the art. As an example, the following protocol (Triton X-100 lysis, phenol/SDS RNase inactivation) gives excellent results with spleen and hybridoma cells (the addition of VRC (veronal ribosyl complex) as an RNase inhibitor is necessary for spleen cells). Guanidinium isothiocyanate/CsCl procedures (yielding total cellular RNA) also give good results but are more time-consuming.

1. Harvest 1 to 5×10^7 cells by centrifugation in a bench top centrifuge at $800 \times g$ for 10 minutes at 4°C . Resuspend gently in 50ml of cold PBS buffer. Centrifuge the cells again at $800 \times g$ for 10 minutes at 4°C , and discard supernatant.

2. On ice, add 1 ml ice-cold lysis buffer to the pellet and resuspend it with a 1ml Gilson pipette by gently pipetting up and down. Leave on ice for 5 minutes.

3. After lysis, remove cell debris by centrifuging at 1300 rpm for 5 minutes in a microfuge at 4°C , in precooled tubes.

4. Transfer 0.5 ml of the supernatant to each of two eppendorfs containing $60 \mu\text{l}$ 10% (w/v) SDS and $250 \mu\text{l}$ phenol (previously equilibrated with 100 mM Tris-HCl pH 8.0). Vortex hard for 2 minutes, then microfuge (13000 rpm) for five minutes at room temperature. Transfer the upper, aqueous, phase to a fresh tube.

5. Re-extract the aqueous upper phase five times with 0.5 ml of phenol.

6. Precipitate with 1/10 volume 3M sodium acetate and 2.5 volumes ethanol at 20°C overnight or dry ice/isopropanol for 30 minutes.

7. Wash the RNA pellet and resuspended in $50 \mu\text{l}$ to check concentration by OD260 and check $2 \mu\text{g}$ on a 1% agarose gel. $40 \mu\text{g}$ of RNA was obtained from spleen cells derived from mice.

Lysis buffer is [10mM Tris-HCl pH 7.4, 1mM MgCl_2 , 150mM NaCl, 10mM VRC (New England Biolabs), 0.5% (w/v) Triton X-100], prepared fresh.

[0290] Lysis buffer is [10mM Tris-HCl pH 7.4, 1mM MgCl_2 , 150mM NaCl, 10mM VRC (New England Biolabs), 0.5% (w/v) Triton X-100], prepared fresh.

B. cDNA Preparation

[0291] cDNA can be prepared using many procedures well known to those skilled in the art. As an example, the following protocol can be used:

1. Set up the following reverse transcription mix:

	μl
H_2O (DEPC-treated)	20
5mM dNTP	10
10 x first strand buffer	10
0.1M DTT	10
FOR primer(s) (10 pmol/ μl)	2 (each) (see below)
RNasin (Promega; 40 U/ μl)	4

NB

i) DEPC is diethylpyrocarbonate, the function of which is to inactivate any enzymes that could degrade DNA

or RNA

ii) dNTP is deoxynucleotide triphosphate

iii) DTT is dithiothreitol the function of which is as an antioxidant to create the reducing environment necessary for enzyme function.

5 iv) RNasin is a ribonuclease inhibitor obtained from Promega Corporation, 2800 Woods Hollow Road, Madison, Wisconsin, USA.

2. Dilute 10 µg RNA to 40 µl final volume with DEPC-treated water. Heat at 65°C for 3 minutes and hold on ice for one minute (to remove secondary structure).

10 3. Add to the RNA the reverse transcription mix (58 µl) and 4 µl of the cloned reverse transcriptase 'Super RT' (Anglian Biotech Ltd., Whitehall House, Whitehall Road, Colchester, Essex) and incubate at 42°C for one hour.

4. Boil the reaction mix for three minutes, cool on ice for one minute and then spin in a microfuge to pellet debris. Transfer the supernatant to a new tube.

15 **[0292]** 10 x first strand buffer is [1.4M KCl, 0.5M Tris-HCl pH 8.1 at 42°C 80mM MgCl₂].

[0293] The primers anneal to the 3' end. Examples of kappa light chain primers are MJK1FONX, MJK2FONX, MJK4FONX and MJK5FONX (provided under 'Primer Sequences' below) and examples of heavy chain primers are MIGG1, 2 (CTG GAC AGG GAT CCA GAG TTC CA) and MIGG3 (CTG GAC AGG GCT CCA TAG TTC CA) which anneal to CH1.

20 **[0294]** Alternatively, any primer that binds to the 3' end of the variable regions VH, VLK, VL, or to the constant regions CH1, CK or CL can be used.

C. Primary PCRs

25 **[0295]** For each PCR and negative control, the following reactions are set up (e.g. one reaction for each of the four VLKs and four VH PCRs). In the following, the Vent DNA polymerase sold by (C.P. Laboratories Ltd (New England Biolabs) address given above) was used. The buffers are as provided by C.P. Laboratories.

	µl
H ₂ O	32.5
10 x Vent buffer	5
20 x Vent BSA	2.5
5mM dNTPs	1.5
FOR primer 10 pmol/µl)	2.5
BACK primer 10pmol/µl	2.5

The FOR and BACK primers are given in the section below entitled 'Primer Sequences'. For VH, the FOR primer is VH1FOR-2 and the BACK primer is VH1BACK. For VLK the FOR primers are MJK1FONX, MJK2FONX, MJK4FONX and MJK5FONX (for the four respective kappa light chains) and the BACK primer is VK2BACK. Only one Kappa light chain BACK primer is necessary, because binding is to a nucleotide sequence common to the four kappa light chains.

40 **[0296]** UV this mix 5 minutes. Add 2.5 µl cDNA preparation (from B above), 2 drops paraffin oil (Sigma Chemicals. Poole, Dorset, UK). Place on a cycling heating block. e.g. PHC-2 manufactured by Techne Ltd. Duxford UK, or set at 94°C. Add 1 µl Vent DNA polymerase under the paraffin. Amplify using 25 cycles of 94°C 1 min, 72°C 2 min. Post-treat at 60°C for 5 min.

45 **[0297]** Purify on a 2% 1mp (low melting point agarose/TAE (tris-acetate EDTA)gel and extract the DNA to 20 µl H₂O per original PCR using a GeneClean kit (see earlier) in accordance with the manufacturers instructions.

D. Preparation of linker

50 **[0298]** Set up in bulk (e.g. 10 times)

	µl
H ₂ O	34.3
10 x Vent buffer	5
20 x Vent BSA	2.5

EP 0 774 511 B1

(continued)

	μl
5mM dNTPs	2
LINKFOR primer 10 pmol/ μl)	2.5
LINKBACK primer 10pmol/ μl	2.5
DNA from scFv D1.3 (example 2)	1
Vent enzyme	0.2

The FOR and BACK primers are given in the section below entitled 'Primer Sequences'. The FOR primer is LINKFOR and the BACK primer is LINKBACK. Cover with paraffin and place on the cycling heating block (see above) at 94°C. Amplify using 25 cycles of 94°C 1 min, 65°C 1 min, 72°C 2 min. Post-treat at 60°C for 5 min.

[0299] Purify on 2% 1mp/TAE gel (using a loading dye without bromophenol blue as a 93bp fragment is desired) and elute with SPIN-X column (Costar Limited, 205 Broadway, Cambridge, Ma. USA..) and precipitation. Take up in 5 μl H₂O per PCR reaction.

E. Assembly PCRs

[0300] A quarter of each PCR reaction product (5 μl) is used for each assembly. The total volume is 25 μl .

[0301] For each of the four VLK primers, the following are set uD:

H ₂ O	4.95
10 x Vent buffer	2.5
20 x Vent BSA	1.25
5mM dNTPs	0.8

UV irradiate this mix for 5 min. Add 5 μl each of Vh and VK band from the primary PCRs and 1.5 μl of linker as isolated from the preparative gels and extracted using the GeneClean kit as described in C and D above. Cover with paraffin. Place on the cycling heating block preset at 94°C. Add 1ul Vent under the paraffin. Amplify using 7 cycles of 94°C 2 min, 72°C 4 min. Then return the temperature to 94°C.

[0302] Add 1.5ul each of VH1BACK and the appropriate VKFOR primers MJK1FONX, MJK2FONX, MJK4FONX or MJK5FONX (10 pmol/ μl) at 94°C. The primers should have been UV-treated as above. Amplify using 20 cycles of 94°C 1.5 min, 72°C 2.5 min. Post-treat at 60°C for 5 min. Purify on 2% 1mp/TAE gel and extract the DNA to 20 μl H₂O per assembly PCR using a GeneClean kit (see earlier) in accordance with the manufacturers instructions.

F. Adding Restriction Sites

[0303] For each assembly and control set up:

	μl
H ₂ O	36.5
10 x Taq buffer	5
5mM dNTPs	2
FOR primer (10 pmol/ μl)	2.5
BACK primer (10 pmol/ μl)	2.5
Assembly product	1

The FOR and BACK primers are given in the section below entitled 'Primer Sequences'. The FOR primer is any of JK1NOT10, JK2NOT10, JK4NOT10 or JK5NOT10 (for the four respective kappa light chains) for putting a Not1 restriction site at the VLK end. The BACK primer is HBKAPA10 for putting an ApaL1 restriction site at the VH end.

[0304] Cover with paraffin and place on the cycling heating block preset at 94°C. Add 0.5 μl Cetus Taq DNA polymerase (Cetus/perkin-Elmer, Beaconsfield, Bucks, UK) under the paraffin. Amplification is carried out using 11 to 15 rounds of cycling (depends on efficiency) at 94°C 1 min, 55°C 1 min, 72°C 2 min. Post-treat at 60°C for 5 min.

[0305] 10 x Taq buffer is [0.1M Tris-HCl pH 8.3 at 25°C, 0.5M KCl, 15mM MgCl₂, 1mg/ml gelatin].

G. Work-up

[0306] Purify once with CHCl_3/IAA (isoamylalcohol), once with phenol, once with CHCl_3/IAA and back-extract everything to ensure minimal losses. Precipitate and wash twice in 70% EtOH. Dissolve in 70 μl H_2O .

Digest overnight at 37°C with NotI:	μl
DNA (joined seq)	70
NEB NotI buffer x 10	10
NEB BSA x 10	10
NotI (10 U/ μl)	10

The DNA (joined sequence) above refers to the assembled DNA sequence comprising in the 5' to 3' direction

ApaL1 restriction site
VH sequence
Linker sequence
VLK sequence
Not 1 restriction site.

[0307] The VLK sequence may be any one of four possible kappa chain sequences.

[0308] The enzymes Not 1 above, ApaL1 below and the buffers NEB Not 1, NEB BSA above and the NEB buffer 4 (below) are obtainable from CP Laboratories, New England Biolabs mentioned above.

[0309] Re-precipitate, take up in 80 μl H_2O . Add to this 10 μl NEB buffer 4 and 10 μl ApaL1.

[0310] Add the enzyme ApaL1 in aliquots throughout the day, as it has a short half-life at 37°C.

[0311] Purify on 2% 1mp/TAE gel and extract the DNA using a GeneClean kit, in accordance with the manufacturers instructions. Redigest if desired.

H. Final DNA product

[0312] The final DNA product is an approximate 700 bp fragment with Apa L1 and Not1 compatible ends consisting of randomly associated heavy and light chain sequences linked by a linker. A typical molecule of this type is the scFvD1.3 molecule incorporated into fdscFvD1.3 described in example 3. These molecules can then be ligated into suitable fd derived vectors, e.g. fdCAT2 (example 5), using standard techniques.

Primer sequences

Primary PCR oligos (restrictions sites underlined):

VH1FOR-2 TGA GGA GAC GGT GAC CGT GGT CCC TTG GCC CC
VH1BACK AGG TSM ARC TGC AGS AGT CWG G
MJK1FONX CCG TTT GAT TTC CAG CTT GGT GCC
MJK2FONX CCG TTT TAT TTC CAG CTT GGT CCC
MJK4FONX CCG TTT TAT TTC CAA CTT TGT CCC
MJK5FONX CCG TTT CAG CTC CAG CTT GGT CCC
VK2BACK GAC ATT GAG CTC ACC CAG TCT CCA

Ambiguity codes M = A or C, R = A or G, S = G or C.

W = A or T

PCR oligos to make linker:

LINKFOR TGG AGA CTC GGT GAG CTC AAT GTC
LINKBACK GGG ACC ACG GTC ACC GTC TCC TCA

For adding restriction sites:

[0313]

```

5      HBKAPA10  CAT GAC CAC AGT GCA CAG GTS MAR CTG CAG SAG TCW
          GG
      JKINOT10  GAG TCA TTC TGC GGC CGC CCG TTT GAT TTC CAG CTT
          GGT GCC
10     JK2NOT10  GAG TCA TTC TGC GGC CGC CCG TTT TAT TTC CAG CTT
          GGT CCC
      JK4NOT10  GAG TCA TTC TGC GGC CGC CCG TTT TAT TTC CAA CTT
          TGT CCC
      JK5NOT10  GAG TCA TTC TGC GGC CGC CCG TTT CAG CTC CAG CTT
15     GGT CCC

```

Example 13. Construction of Phagemid Containing GeneIII fused with the Coding Sequence for a Binding Molecule

[0314] It would be useful to improve the transfection efficiency of the phage-binding molecule system and also to have the possibility of displaying different numbers and specificities of binding molecules on the surface of the same bacteriophage. The applicants have devised a method that achieves both aims.

[0315] The approach is derived from the phagemid system based on pUC119 [Vieira, J and Messing, J. (1987) Methods Enzymol. 153:3]. In brief, gene III from fd-CAT2 (example 5) and gene III scFv fusion from fd-CAT2 scFv D1.3 (example 2) were cloned downstream of the lac promoter in separate samples of pUC119, in order that the inserted gene III and gene III fusion could be 'rescued' by M13M07 helper phage [Vieira, J and Messing, J. et supra.] prepared according to Sambrook et al. 1989 supra. The majority of rescued phage would be expected to contain a genome derived from the pUC119 plasmid that contains the binding molecule-gene III fusion and should express varying numbers of the binding molecule on the surface up to the normal maximum of 3-5 molecules of gene III of the surface of wild type phage. The system has been exemplified below using an antibody as the binding molecule.

[0316] An fdCAT2 containing the single chain Fv form of the D1.3 antilysozyme antibody was formed by digesting fdTscFvD1.3 (example 2) with Pst1 and Xho1, purifying the fragment containing the scFv fragment and ligating this into Pst1 and Xho1 digested fdCAT2. The appropriate clone, called fdCAT2 scFvD1.3 was selected after plating onto 2xTY tetracycline (15µg/ml) and confirmed by restriction enzyme and sequence analysis.

[0317] Gene III from fd-CAT2 (example 5) and the gene III scFv fusion from fd-CAT2 scFvD1.3 was PCR-amplified using the primers A and B shown below:

Primer A: **TGC GAA GCT TTG GAG CCT TTT TTT TTG GAG ATT TTC**
AAC G

Primer B: **CAG TGA ATT CCT ATT AAG ACT CCT TAT TAC GCA GTA**
TGT TAG C

Primer A anneals to the 5' end of gene III including the ribosome binding site is located and incorporates a Hind III site. Primer B anneals to the 3' end of gene III at the C-terminus and incorporates two UAA stop codons and an EcoR1 site. 100 ng of fd-CAT2 and fd-CAT2 scFv D1.3 DNA was used as templates for PCR-amplification in a total reaction volume of 50µl as described in example 7, except that 20 cycles of amplification were performed: 94°C 1 minute, 50°C 1 minute, 72°C 3 minutes. This resulted in amplification of the expected 1.2Kb fragment from fd-CAT2 and a 1.8Kb fragment from fd-CAT2 scFv D1.3.

[0318] The PCR fragments were digested with EcoR1 and Hind III, gel-purified and ligated into Eco-R1- and Hind III-cut and dephosphorylated pUC119 DNA and transformed into E.coli TG1 using standard techniques (Sambrook et al., et supra). Transformed cells were plated on SOB agar (Sambrook et al. 1989 supra) containing 100µg/ml ampicillin and 2% glucose. The resulting clones were called pCAT-3 (derived from fd-CAT2) and pCAT-3 scFv D1.3 (derived from fd-CAT2 scFv D1.3).

Example 14 Rescue of Anti-Lysozyme Antibody Specificity from pCAT-3 scFv D1.3 by M13K07

[0319] Single pCAT-3 and pCAT-3 scFv D1.3 colonies were picked into 1.5ml 2TY containing 100µg/ml ampicillin and 2% glucose, and grown 6 hrs at 30°C. 30µl of these stationary cells were added to 6mls 2YT containing 100µg/ml ampicillin and 2% glucose in 50ml polypropylene tubes (Falcon, Becton Dickinson Labware, 1950 Williams Drive, Oxnard, CA. USA) and grown for 1.5 hrs at 30°C at 380rpm in a New Brunswick Orbital Shaker (New Brunswick Scientific Ltd., Edison House 163 Dixons Hill road, North Mimms, Hatfield, UK). Cells were pelleted by centrifugation at 5,000g for 25 minutes and the tubes drained on tissue paper. The cell pellets were then suspended in 6mls 2TY containing 1.25×10^9 p.f.u. ml⁻¹ M13K07 bacteriophage added. The mixture was left on ice for 5 minutes followed by growth at 35°C for 45 minutes at 450rpm. A cocktail was then added containing 4µl 100µg/ml ampicillin, 0.5µl 0.1M IPTG and 50µl 10mg/ml kanamycin, and the cultures grown overnight at 35°C, 450rpm.

[0320] The following day the cultures were centrifuged and phage particles PEG precipitated as described in example 6. Phage pellets were resuspended in 100µl TE (tris-EDTA see example 6) and phage titred on E.coli TG1. Aliquots of infected cells were plated on 2TY containing either 100µg/ml ampicillin to select for pUC119 phage particles. or 50µg/ml kanamycin to select for the M13 K07 helper phage. Plates were incubated overnight at 37°C and antibiotic-resistant colonies counted:

DNA	amp ^R	kan ^R
pCAT-3	1.8×10^{11} colonies	1.2×10^9 colonies
pCAT-3scFv D1.3	2.4×10^{11} colonies	2.0×10^9 colonies

This shows that the amp^R phagemid particles are infective and present in the rescued phage population at a 100-fold excess over kan^R M13K07 helper phage.

[0321] Phage were assayed for anti-lysozyme activity by ELISA as described in example 6, with the following modifications:

- 1) ELISA plates were blocked for 3 hrs with 2% Marvel/PBS.
- 2) 50µl phage, 400µl 1xPBS and 50µl 20% Marvel were mixed end over end for 20 minutes at room temperature before adding 150µl per well.
- 3) Phage were left to bind for 2 hours at room temperature.
- 4) All washes post phage binding were:
 - 2 quick rinses PBS/0.5% Tween 20
 - 3x2 minute washes PBS/0.5% Tween 20
 - 2 quick rinses PBS no detergent
 - 3x2 minute washes PBS no detergent

[0322] The result of this ELISA is shown in figure 19, which shows that the antibody specificity can indeed be rescued efficiently.

[0323] It is considered a truism of bacterial genetics that when mutant and wild-type proteins are co-expressed in the same cell, the wild-type protein is used preferentially. This is analogous to the above situation wherein mutant (i. e. antibody fusion) and wild-type gene III proteins (from M13K07) are competing for assembly as part of the pUC119 phagemid particle. It is therefore envisaged that the majority of the resulting pUC 119 phage particles will have fewer gene III-antibody fusion molecules on their surface than is the case for purely phage system described for instance in example 2. Such phagemid antibodies are therefore likely to bind antigen with a lower avidity than fd phage antibodies with three or more copies of the antibody fusion on their surfaces (there is no wild-type gene III, in the system described, for instance, in example 2), and provide a route to production of phage particles with different numbers of the same binding molecule (and hence different acidities for the ligand/antigen) or multiple different binding specificities on their surface, by using helper phage such as M13K07 to rescue cells expressing two or more gene III-antibody fusions.

[0324] It is also possible to derive helper phage that do not encode a functional gene III in their genomes (by for example deleting the gene III sequence or a portion of it or by incorporating an amber mutation within the gene). These defective phages will only grow on appropriate cells (for example that provide functional gene III in trans, or contain an amber suppressor gene), but when used to rescue phage antibodies, will only incorporate the gene III antibody fusion encoded by the phagemid into the released phage particle.

Example 15. Transformation Efficiency of pCAT-3 and pCAT-3 scFv D1.3 phagemids

[0325] pUC 19, pCAT-3 and pCAT-3 scFv D1.3 plasmid DNAs, and fdCAT-2 phage DNA was prepared, and used to transform E.coli TG1, pCAT-3 and pCAT-3 scFv D1.3 transformations were plated on SOB agar containing 100µg/ml ampicillin and 2% glucose, and incubated overnight at 30°C. fdCAT-2 transformations were plated on TY agar containing 15µg/ml tetracycline and incubated overnight at 37°C. Transformation efficiencies are expressed as colonies per µg of input DNA.

DNA	Transformation efficiency
pUC 19	1.10^9
pCAT-3	1.10^8
pCAT-3scFv D1.3	1.10^8
fd CAT-2	8.10^5

[0326] As expected, transformation of the phagemid vector is approximately 100-fold more efficient than the parental fdCAT-2 vector. Furthermore, the presence of a scFv antibody fragment does not compromise efficiency. This improvement in transformation efficiency is practically useful in the generation of phage antibodies libraries that have large repertoires of different binding specificities.

Example 16PCR Assembly of a Single Chain Fv Library from an Immunised Mouse

[0327] To demonstrate the utility of phage for the selection of antibodies from repertoires, the first requirement is to be able to prepare a diverse, representative library of the antibody repertoire of an animal and display this repertoire on the surface of bacteriophage fd.

[0328] Cytoplasmic RNA was isolated according to example 12 from the pooled spleens of five male Balb/c mice boosted 8 weeks after primary immunisation with 2-phenyl-5-oxazolone (ph OX) coupled to chicken serum albumin. cDNA preparation and PCR assembly of the mouse VH and VL kappa repertoires for phage display was as described in example 12. The molecules thus obtained were ligated into fdCAT2.

[0329] Vector fdCAT2 was extensively digested with Not1 and Apa1., purified by electroelution (Sambrook et al. 1989 supra) and 1 µg ligated to 0.5 µg (5 µg for the hierarchical libraries: see example 18) of the assembled scFv genes in 1 ml with 8000 units T4 DNA ligase (New England Biolabs). The ligation was carried out overnight at 16°C. Purified ligation mix was electroporated in six aliquots into MC1061 cells (W. J. Dower, J. F. Miller & C. W. Ragsdale Nucleic Acids Res. 16 6127-6145 1988) and plated on NZY medium (Sambrook et al. 1989 supra) with 15µg/ml tetracycline, in 243x243 mm dishes (Nunc): 90-95% of clones contained scFv genes by PCR screening.

Recombinant colonies were screened by PCR (conditions as in example 7 using primers VH1BACK and MJK1FONX, MJK2FONX, MJK4FONX and MJK5FONX (see example 12) followed by digestion with the frequent cutting enzyme BstN1 (New England Biolabs, used according to the manufacturers instructions). The library of 2×10^5 clones appeared diverse as judged by the variety of digestion patterns seen in Figure 20, and sequencing revealed the presence of most VH groups (R. Dildrop, Immunol. Today 5 85-86. 1984) and VK subgroups (Kabat. E.A. et al. 1987 supra) (data not shown). None of the 568 clones tested bound to phOx as detected by ELISA as in example 9.

[0330] Thus the ability to select antibody provided by the use of phage antibodies (as in example 17) is essential to readily isolate antibodies with antigen binding activity from randomly combined VH and VL domains. Very extensive screening would be required to isolate antigenbinding fragments if the random combinatorial approach of Huse et al. 1989 (supra) were used.

Example 17Selection of Antibodies Specific for 2-phenyl-5-oxazolone from a Repertoire Derived from an Immunised Mouse

[0331] The library prepared in example 16 was used to demonstrate that ability of the phage system to select antibodies on the basis of their antibody specificity.

[0332] None of the 568 clones tested from the unselected library bound to phOx as detected by ELISA.

[0333] Screening for binding of the phage to hapten was carried out by ELISA: 96-well plates were coated with 10 µg/ml phOx-BSA or 10 µg/ml BSA in phosphate-buffered saline (PBS) overnight at room temperature. Colonies of

phage-transduced bacteria were inoculated into 200 μ l 2 x TY with 12.5 μ g/ml tetracycline in 96-well plates ('cell wells', Nuclon) and grown with shaking (300 rpm) for 24 hours at 37°C. At this stage cultures were saturated and phage titres were reproducible (10^{10} TU/ml). 50 μ l phage supernatant, mixed with 50 μ l PBS containing 4% skimmed milk powder, was then added to the coated plates. Further details as in example 9.

5 **[0334]** The library of phages was passed down a phOx affinity column (Table 3A), and eluted with hapten. Colonies from the library prepared in example 18 were scraped into 50ml 2 x TY medium and shaken at 37°C for 30 min. Liberated phage were precipitated twice with polyethylene glycol and resuspended to 10^{12} TU (transducing units)/ml in water (titred as in example 8). For affinity selection, a 1 ml column of phOx-BSA-Sepharose (O. Makela, M. Kaartinen, J.L. T. Pelonen and K. Karjalainen J. Exp. Med. 148 1644-1660, 1978) was washed with 300 ml phosphate-buffered saline (PBS), and 20 ml PBS containing 2% skimmed milk powder (MPBS). 10^{12} TU phage were loaded in 10 ml MPBS, washed with 10 ml MPBS and finally 200 ml PBS. The bound phage were eluted with 5 ml 1 mM 4- ϵ -amino-caproic acid methylene 2-phenyloxazol-5-one (phOx-CAP; O. Makela et al. 1978, supra). About 10^6 TU eluted phage were amplified by infecting 1 ml log phase E.coli TG1 and plating as above. For a further round of selection, colonies were scraped into 10 ml 2 x TY medium and then processed as above. Of the eluted clones, 13% were found to bind to phOx after the first round selection, and ranged from poor to strong binding in ELISA.

15 **[0335]** To sequence clones, template DNA was prepared from the supernatants of 10 ml cultures grown for 24 hours, and sequenced using the dideoxy method and a Sequenase kit (USB), with primer LINKFOR (see example 12) for the VH genes and primer fdSEQ1 (5'-GAA TTT TCT GTA TGA GG) for the Vk genes. Twenty-three of these hapten-binding clones were sequenced and eight different VH genes (A to H) were found in a variety of pairings with seven different
20 Vk genes (a to g) (Fig. 21). Most of the domains, such as VH-B and Vk-d were 'promiscuous', able to bind hapten with any of several partners.

[0336] The sequences of the V-genes were related to those seen in the secondary response to phOx, but with differences (Fig. 21). Thus phOx hybridomas from the secondary response employ somatically mutated derivatives of three types of Vk genes - Vkoxl, 'Vkox-like' and Vk45.1 genes (C. Berek, G. M. Griffiths & C. Milstein Nature 316
25 412-418 (1985). These can pair with VH genes from several groups, from Vkoxl more commonly pairs with the VHoxl gene (VH group 2. R.Dildrop supra). Vkoxl genes are always, and Vkox-like genes often, found in association with heavy chains (including VHoxl) and contain a short five residue CDR3, with the sequence motif Asp-X-Gly-X-X in which the central glycine is needed to create a cavity for phOx. In the random combinatorial library however, nearly all of the VH genes belonged to group 1, and most of the Vk genes were ox-like and associated with VH domains with a five
30 residue CDR3, motif Asp/Asn-X-Gly-X-X (Fig. 21). Vkoxl and VHoxl were found only once (Vk-f and VH-E), and not in combination with each other. Indeed Vk-f lacks the Trp91 involved in phOx binding and was paired with a VH (VH-C) with a six residue CDR3.

[0337] A matrix combination of VH and VK genes was identified in phOx-binding clones selected from this random combinatorial library. The number of clones found with each combination are shown in Fig. 22. The binding to phOx-BSA, as judged by the ELISA signal, appeared to vary (marked by shading in Fig. 22). No binding was seen to
35 BSA alone.

[0338] A second round of selection of the original, random combinatorial library from immune mice resulted in 93% of eluted clones binding phOx (Table 3). Most of these clones were Vk-d combinations, and bound strongly to phOx in ELISA (data not shown). Few weak binders were seen. This suggested that affinity chromatography had not only
40 enriched for binders, but also for the best.

[0339] Florescence quench titrations determined the Kd of VH-B/Vk-d for phOx-GABA as 10^{-8} M (example 19), indicating that antibodies with affinities representative of the secondary response can be selected from secondary response, only two (out of eleven characterised) secrete antibodies of a higher affinity than VH-B/Vk-d (C. Berek et al. 1985 supra). The Kd of VH-B/Vk-b for phOx-GABA was determined as 10^{-5} M (example 19). Thus phage bearing scFv fragments with weak affinities can be selected with antigen, probably due to the avidity of the multiple antibody heads
45 on the phage.

[0340] This example shows that antigen specificities can be isolated from libraries derived from immunised mice. It will often be desired to express these antibodies in a soluble form for further study and for use in therapeutic and diagnostic applications. Example 19 demonstrates determination of the affinity of soluble scFv fragments selected using phage antibodies. Example 23 demonstrates that soluble fragments have similar properties to those displayed
50 on phage. For many purposes it will be desired to construct and express an antibody molecule which contains the Fc portions of the heavy chain, and perhaps vary the immunoglobulin isotype. To accomplish this, it is necessary to sub-clone the antigen binding sites identified using the phage selection system into a vector for expression in mammalian cells, using methodology similar to that described by Orlandi, R. et al. (1989, supra). For instance, the VH and VL genes could be amplified separately by PCR with primers containing appropriate restriction sites and inserted into
55 vectors such as pSV-gpt HuIgG1 (L. Riechmann et al Nature 332 323-327, 1988) which allows expression of the VH domain as part of a heavy chain IgG1 isotype and pSV-hyg HuCK which allows expression of the VL domain attached to the K light chain constant region. Furthermore, fusions of VH and VL domains can be made with genes encoding

non-immunoglobulin proteins, for example, enzymes.

Example 18

5 Generation of Further Antibody Specificities by the Assembly of Hierarchical Libraries

[0341] Further antibody specificities were derived from the library prepared and screened in examples 16 and 17 using a hierarchical approach.

[0342] The promiscuity of the VH-B and Vk-d domains prompted the applicants to force further pairings, by assembling these genes with the entire repertoires of either Vk or VH genes from the same immunised mice. The resulting 'hierarchical' libraries, (VH-B x Vk-rep and VH-rep x Vk-d), each with 4×10^7 members, were subjected to a round of selection and hapten-binding clones isolated (Table 3). As shown by ELISA, most were strong binders. By sequencing twenty-four clones from each library, the applicants identified fourteen new partners for VH-B and thirteen for Vk-d (Fig. 21). Apart from VH-B and Vk-c, none of the previous partners (or indeed other clones) from the random combinatorial library was isolated again. Again the Vk genes were mainly ox-like and the VH genes mainly group 1 (as defined in Dildrop, R. 1984 *supra*), but the only examples of Vkoxl (Vk-h, -p, -q and -r) have Trp91, and the VH-CDR3 motif Asp-X-Gly-X-X now predominates. Thus some features of the phOx hybridomas seemed to emerge more strongly in the hierarchical library. The new partners differed from each other mainly by small alterations in the CDRs, indicating that much of the subtle diversity had remained untapped by the random combinatorial approach. More generally it has been shown that a spectrum of related antibodies can be made by keeping one of the partners fixed and varying the other, and this could prove invaluable for fine tuning of antibody affinity and specificity.

[0343] Therefore, again, phage antibodies allow a greater range of antibody molecules to be analysed for desired properties.

[0344] This example, and example 17, demonstrate the isolation of individual antibody specificities through display on the surface of phage. However, for some purposes it may be more desirable to have a mixture of antibodies, equivalent to a polyclonal antiserum (for instance, for immunoprecipitation). To prepare a mixture of antibodies, one could mix clones and express soluble antibodies or antibody fragments or alternatively select clones from a library to give a highly enriched pool of genes encoding antibodies or antibody fragments directed against a ligand of interest and express antibodies from these clones.

30 Example 19

35 Selection of Antibodies Displayed on Bacteriophage with Different Affinities for 2-phenyl-5-oxazolone using Affinity Chromatography

[0345] The ELISA data shown in example 17 suggested that affinity chromatography had not only enriched for binders, but also for the best. To confirm this, the binding affinities of a strong binding and a weak binding phage were determined and then demonstrated that they could be separated from each other using affinity chromatography.

[0346] Clones VH-B/Vk-b and VH-B/Vk-d were reamplified with MJK1FONX, MJK2FONX, MJK4FONX and MJK5FONX (see example 12) and VH1BACK-Sfil (5'-TCG CCG CCC AGC, CCG CCA TGG CC(G/C) AGG T(C/G) (A/C) A(A/G)C TGC AG(C/G) AGT C(A/T)G G), a primer that introduces an Sfil site (underlined) at the 5' end of the VH gene. VH-B/Vk-d was cloned into a phagemid e.g. pJM1 (a gift from A. Griffiths and J. Marks) as an Sfil-NotI cassette, downstream of the pelB leader for periplasmic secretion (M. Better *et al.* *supra*), with a C-terminal peptide tag for detection (see example 20 and figure), and under the control of a P_L promoter (H. Shimatake & M. Rosenberg Nature 292 128-132 1981). The phagemid should have the following features: a) unique Sfil and NotI restriction sites downstream of a pelB leader; b) a sequence encoding a C-terminal peptide tag for detection; and c) a λ P_L promoter controlling expression. 10 litre cultures of E.coli N4830-1 (M. E. Gottesman, S. Adhya & A. Das J.Mol.Biol 140 57-75 1980) harbouring each phagemid were induced as in K. Nagai & H. C. Thogerson (Methods Enzymol 153 461-481 1987) and supernatants precipitated with 50% ammonium sulphate. The resuspended precipitate was dialysed into PBS + 0.2 mM EDTA (PBSE), loaded onto a 1.5ml column of phOx-Sepharose and the column washed sequentially with 100 ml PBS: 100 ml 0.1 M Tris-HCl, 0.5 M NaCl, pH 8.0: 10ml 50 mM citrate, pH 5.0: 10 ml 50 mM citrate, pH4.0, and 20 ml 50 mM glycine, pH 3.0. scFv fragments were eluted with 50 mM glycine, pH 2.0, neutralised with Tris base and dialysed against PBSE. VH-B/Vk-b was cloned into a phagemid vector based on pUC119 encoding identical signal and tag sequences to pJM1, and expression induced at 30°C in a 10 litre culture of E.coli TG1 harbouring the phagemid as in D. de Bellis & I. Schwartz (1980 Nucleic Acids Res 18 1311). The low affinity of clone VH-B/Vk-b made its purification on phOx-Sepharose impossible. Therefore after concentration by ultrafiltration (Filtron, Flowgen), the supernatant (100 ml of 600 ml) was loaded onto a 1 ml column of protein A-Sepharose coupled (E. Harlow & D. Lane 1988 *supra*) to the monoclonal antibody 9E10 (Evan, G. I. *et al.* Mol.Cell Biol. 5 3610-3616 1985) that recognises the peptide

tag. The column was washed with 200 ml PBS and 50 ml PBS made 0.5 M in NaCl. scFv fragments were eluted with 100 ml 0.2M glycine, pH 3.0, with neutralisation and dialysis as before.

[0347] The K_d ($1.0 \pm 0.2 \times 10^{-8}$ M) for clone VH-B/Vk-d was determined by fluorescence quench titration with 4-E-amino-butyric acid methylene 2-phenyl-oxazol-5-one (phOxGABA Co. Makela *et al*, 1978 *supra*). Excitation was at 280 nm, emission was monitored at 340 nm and the K_d calculated. The K_d of the low affinity clone VH-B/Vk-b was determined as $1.8 \pm 0.3 \times 10^{-5}$ M (not shown). To minimise light adsorption by the higher concentrations of phOx-GABA required, excitation was at 260 nm and emission was monitored at 304 nm. In addition the fluorescence values were divided by those from a parallel titration of the lysozyme binding Fv fragment D1.3. The value was calculated as in H. N. Eisen *Meth.Med.Res.* 10 115-121 1964. A mixture of clones VH-B/Vk-b and VH-B/Vk-d, 7×10^{10} TU phage in the ratio 20 VH-B/Vk-b : 1 VH-B/Vk-d were loaded onto a phOx-BSA-Sepharose column in 10 ml MPBS and eluted as above. Eluted phage were used to reinfect E.coli TG1, and phage produced and harvested as before. Approximately 10^{11} TU phage were loaded onto a second affinity column and the process repeated to give a total of three column passes. Dilutions of eluted phage at each stage were plated in duplicate and probed separately with oligonucleotides specific for Vk-b (5'-GAG CGG GTA ACC ACT GTA CT) or Vk-d (5'-GAA TGG TAT AGT ACT ACC CT). After these two rounds, essentially all the eluted phage were VH-B/Vk-d (table 3). Therefore phage antibodies can be selected on the basis of the antigen affinity of the antibody displayed.

Example 20

20 Construction of Phagemid pHEN1 for the Expression of Antibody Fragments Expressed on the Surface of Bacteriophage following Superinfection

[0348] The phagemid pHEN1 (figure 23) is a derivative of pUC119 (Vieira, J. & Messing, J. *Methods Enzymol* 153 pp 3-11, 1987). The coding region of g3p from fdCAT2, including signal peptide and cloning sites, was amplified by PCR, using primers G3FUFO and G3FUBA (given below) (which contain EcoRI. and HindIII sites respectively), and cloned as a HindIII-EcoRI fragment into pUC119. The HindIII-NotI fragment encoding the g3p signal sequence was the replaced by a pelB signal peptide (Better, M. *et al.* *Science* 240 1041-1043, 1988) with an internal SfiI site, allowing antibody genes to be cloned as fli-NotI fragments. A peptide tag, c-myc, (Munro, S. & Pelham, H. *Cell* 46 291-300, 1986) was introduced directly after the NotI site by cloning an oligonucleotide cassette, and followed by an amber codon introduced by site-directed mutagenesis using an in vitro mutagenesis kit (Amersham International) (figure 23b).

35 G3FUFO, 5'-CAG TGA ATT CTT ATT AAG ACT CCT TAT TAC GCA GTA
TGT TAG C;
G3FUBA, 5'-TGC GAA GCT TTG GAG CCT TTT TTT TTG GAG ATT TTC
AAC G;

Example 21

40 Display of Single Chain Fv and Fab Fragments Derived from the Anti-Oxazolone Antibody NQ10.12.5 on Bacteriophage fd using pHEN1 and fdCAT2

[0349] A range of constructs (see figure 24) were made from a clone (essentially construct II in pUC19) designed for expression in bacteria of a soluble Fab fragment (Better *et al.* 1988 see above) from the mouse anti-phOx (2-phenyl-5-oxazolone) antibody NQ10.12.5 (Griffiths, G. M. *et al.* *Nature* 312, 271-275, 1984). In construct II, the V-regions are derived from NQ10.12.5 and attached to human Ck and CH1 (γ 1 isotype) constant domains. The C-terminal cysteine residues, which normally form a covalent link between light and heavy antibody chains, have been deleted from both the constant domains. To clone heavy and light chain genes together as Fab fragments (construct II) or as separate chains (constructs III and IV) for phage display, DNA was amplified from construct II by PCR to introduce a NotI restriction site at the 3' end, and at the 5' end either an ApaI site (for cloning into fd-CAT2) or SfiI site (for cloning into pHEN1). The primers FABNOTFOK with VH1BACKAPA (or VH1BACKSFI15) were used for PCR amplification of genes encoding Fab fragments (construct II), the primers FABNOTFOH with VH1BACKAPA (or VH1BACKSFI15) for heavy chains (construct III), and the primers FABNOTFOK and MVKBASFI (or MVKBASFI) for light chains (construct IV).

[0350] The single-chain Fv version of NQ10.12.5 (construct I) has the heavy (VH) and light chain (Vk) variable domains joined by a flexible linker (Gly₄Ser)₃ (Huston, J. S. *et al.* *Proc. Natl. Acad. Sci. USA* 85 5879-5883, 1988) and was constructed from construct II by 'splicing by overlap extension' as in example 12. The assembled genes were reamplified with primers VK3F2NOT and VH1BACKAPA (or VH1BACKSFI15) to append restriction sites for cloning

into fd-CAT2 (ApaLI-NotI) or pHEN1 (SfiI-NotI).

```

5  VH1BACKAPA, 5'-CAT GAC CAC AGT GCA CAG GT(C/G) (A/C)A(A/G)
   CTG CAG (C/G)AG TC(A/T) GG;
   VH1BACKSFI15, 5'-CAT GCC ATG ACT CGC GGC CCA GCC GGC CAT
   GGC C(C/G)A GGT (C/G)(A/C)A (A/G)CT GCA G(C/G)A GTC
   (A/T)GG;
   FABNOTFOH, 5'-CCA CGA TTC TGC GGC CGC TGA AGA TTT GGG CTC
   AAC TTT CTT GTC GAC;
10  FABNOTFOK, 5'-CCA CGA TTC TGC GGC CGC TGA CTC TCC GCG GTT
   GAA GCT CTT TGT GAC;
   MVKBAAPA, 5'-CAC AGT GCA CTC GAC ATT GAG CTC ACC CAG TCT
   CCA;
15  MVKBASFI, 5'-CAT GAC CAC GCG GCC CAG CCG GCC ATG GCC GAC

   ATT GAG CTC ACC CAG TCT CCA;
20  VK3F2NOT, 5'-TTC TGC GGC CGC CCG TTT CAG CTC GAG CTT GGT
   CCC.

```

Restriction sites are underlined.

25 Rescue of Phage and Phagemid particles

[0351] Constructs I-IV (figure 24) were introduced into both fd-CAT2 and pHEN1. Phage fd-CAT2 (and fd-CAT2-1, II, III or IV) was taken from the supernatant of infected E.coli TG1 after shaking at 37°C overnight in 2xTY medium with 12.5µg/ml tetracycline, and used directly in ELISA. Phagemid pHEN1 (and pHEN1-I and II) in E.coli TG1 (supE) were grown overnight in 2 ml 2xTY medium, 100 µg/ml ampicillin, and 1% glucose (without glucose; expression of g3p prevents later superinfection by helper phage). 10µl of the overnight culture was used to inoculate 2 ml of 2xTY medium, 100µg/ml ampicillin, 1% glucose, and shaken at 37°C for 1 hour. The cells were washed and resuspended in 2xTY, 100 µg/ml ampicillin, and phagemid particles rescued by adding 2 µl (10⁸pfu) VCSM13 helper phage (Stratagene). After growth for one hour, 4µl kanamycin (25 mg/ml) was added, and the culture grown overnight. The phagemid particles were concentrated 10-fold for ELISA by precipitation with polyethylene glycol.

ELISA

[0352] Detection of phage binding to 2-phenyl-5-oxazolone (phOx) was performed as in example 9. 96-well plates were coated with 10 µg/ml phOx-BSA or 10 µg/ml BSA in PBS overnight at room temperature, and blocked with PBSS containing 2% skimmed milk powder. Phage (mid) supernatant (50 µl) mixed with 50 µl PBS containing 4% skimmed milk powder was added to the wells and assayed. To detect binding of soluble scFv or Fab fragments secreted from pHEN1, the c-myc peptide tag described by Munro and Pelham 1986 supra, was detected using the anti-myc monoclonal 9E10 (Evan, G. I. et al. Mol Cell Biol 5 3610-3616, 1985) followed by detection with peroxidase-conjugated goat anti-mouse immunoglobulin. Other details are as in example 9.

[0353] The constructs in fdCAT2 and pHEN1 display antibody fragments of the surface of filamentous phage. The phage vector, fd-CAT2 (figure 8) is based on the vector fd-tet (Zacher, A. N. et al. Gene 9 127-140, 1980) and has restriction sites (ApaLI and NotI) for cloning antibody genes (or other protein) genes for expression as fusions to the N-terminus of the phage coat protein g3p. Transcription of the antibody-g3p fusions in fd-CAT2 is driven from the gene III promoter and the fusion protein targetted to the periplasm by means of the g3p leader. Fab and scFv fragments of NQ10.12.5 cloned into fd-CAT2 for display were shown to bind to phOx-BSA (but not BSA) by ELISA (table 4). Phage were considered to be binding if A₄₀₅ of the sample was at least 10-fold greater than the background in ELISA.

[0354] The phagemid vector, pHEN1 (fig. 23), is based upon pUC119 and contains restriction sites (SfiI and NotI) for cloning the fusion proteins. Here the transcription of antibody-g3p fusions is driven from the inducible lacZ promoter and the fusion protein targetted to the periplasm by means of the pelB leader. Phagemid was rescued with VCSM13 helper phage in 2xTY medium containing no glucose or IPTG: under these conditions there is sufficient expression of antibody-g3p. Fab and scFv fragments of NQ10.12.5 cloned into pHEN1 for display were shown to bind to phOx-BSA (but not BSA) by ELISA (Table 4) using the same criterion as above.

[0355] An alternative methodology for preparing libraries of Fab fragments expressed on the surface of phage would be to:

1. Prepare a library of phage expressing heavy chain (VHCH) genes from inserts in the phage genome.
2. Prepare a library of light chain genes in a plasmid expression vector in E.coli, preferably a phagemid, and isolate the soluble protein light chains expressed from this library.
3. Bind the soluble protein light chains from the library to the heavy chain library displayed on phage.
4. Select phage with the desired properties of affinity and specificity. These will encode the heavy chain (VHCH) genes.
5. Isolate the light chain genes encoding light chains which form suitable antigen binding sites in combination with the selected heavy chains, preferably by using superinfection of bacteria, containing phagemid expressing the light chain, with phage expressing the selected heavy chain (as described in example 20) and then assaying for antigen binding.

15 Example 22

Rescue of Phagemid Encoding a Gene III Protein Fusion with Antibody Heavy or Light Chains by Phage Encoding the Complementary Antibody Chain Displayed on Phage and the Use of this Technique to Make Dual Combinatorial Libraries

[0356] With random combinatorial libraries there is a limitation on the potential diversity of displayed Fab fragments due to the transformation efficiency of bacterial cells. Described here is a strategy (dual combinatorial libraries) to overcome this problem, potentially increasing the number of phage surveyed by a factor of 10^7 .

[0357] For assembly of heavy and light chains expressed from different vectors, phagemid (pHEN1-III or IV) was grown in E.coli HB2151 (a non-suppressor strain) to allow production of soluble chains, and rescued as above (example 23) except that helper phage were used expressing partner chains as fusions to g3p (10^9 TU fd-CAT2-IV or III respectively) and 2 μ l tetracycline (12.5 mg/ml) in place of kanamycin.

30 **Separate Vectors to Encode Fab Heavy and Light Chains**

[0358] The heavy and light chains of Fab fragments can be encoded together in the same vector (example 21) or in different vectors. To demonstrate this the heavy chain (construct III) was cloned into pHEN1 (to provide soluble fragments) and the light chain (construct IV) into fd-CAT2 (to make the fusion with g3p). The phagemid pHEN1-III, grown in E.coli HB2151 (non-suppressor) was rescued with fd-CAT2-IV phage, and phage(mid) shown to bind to pHx:BSA, but not to BSA (Table 4). This demonstrates that soluble light chain is correctly associating with the heavy chain anchored to the g3p, since neither heavy chain nor light chain alone bind antigen (Table 4).

[0359] Similar results were obtained in the reverse experiment (with phagemid pHEN1-IV and fd-CAT2-III phage) in which the heavy chain was produced as a soluble molecule and the light chain anchored to g3p (Table 4). Hence a Fab fragment is assembled on the surface of phage by fusion of either heavy or light chain to g3p, provided the other chain is secreted using the same or another vector (figure 25).

[0360] The resulting phage population is a mixture of phage and rescued phagemid. The ratio of the two types of particle was assessed by infecting log phase E.coli TG1 and plating on TYE plates with either 15 μ g/ml tetracycline (to select for fd-CAT2) or 100 μ g/ml ampicillin (to select for pHEN1). The titre of fd-CAT2 phage was 5×10^{11} TU/ml and the titre of pHEN1 2×10^{10} TU/ml, indicating a packaging ratio of 25 phage per phagemid.

[0361] Demonstrated here is an alternative strategy involving display of the heterodimeric antibody Fab fragments on the surface of phage. One of the chains is fused to g3p and the other is secreted in soluble form into the periplasmic space of the E.coli where it associates non-covalently with the g3p fusion, and binds specifically to antigen. Either the light or heavy chain can be fused to the g3p: they are displayed on the phage as Fab fragments and bind antigen (Figure 25). Described are both phage and phagemid vectors for surface display. Phagemids are probably superior to phage vectors for creation of large phage display libraries. Particularly in view of their higher transfection efficiencies (Two to three orders of magnitude higher), allowing larger libraries to be constructed. The phagemid vector, pHEN1 also allows the expression of soluble Fab fragments in non-suppressor E.coli.

[0362] Also demonstrated here is that heavy and light chains encoded on the same vector (construct II), or on different vectors (constructs III and IV) can be displayed as Fab fragments. This offers two distinct ways of making random combinatorial libraries for display. Libraries of heavy and light chain genes, amplified by PCR, could be randomly linked by a 'PCR assembly' process (example 12) based on 'splicing by overlap extension', cloned into phage(mid) display vectors and expressed from the same promoter as part of the same transcript (construct II) as above, or indeed from different promoters as separate transcripts. Here the phage(mid) vector encodes and displays both chains. For a com-

binatorial library of 10^7 heavy chains and 10^7 light chains, the potential diversity of displayed Fab fragments (10^{14}) is limited by the transfection efficiency of bacterial cells by the vector (about 10^9 clones per μg cut and ligated plasmid at best) (W.J. Dower et al Nucl. Acids. Res. 16 6127-6145, 1988). Libraries thus prepared are analogous to the random combinatorial library method described by Huse, W.D. et al Science 246 1275-1281 (1989), but have the important additional feature that display on the surface of phage gives a powerful method of selecting antibody specificities from the large number of clones generated.

[0363] Alternatively, libraries of heavy and light chains could be cloned into different vectors for expression in the same cell, with a phage vector encoding the g3p fusion and a phagemid encoding the soluble chain. The phage acts as a helper, and the infected bacteria produced both packaged phage and phagemid. Each phage or phagemid displays both chains but encodes only one chain and thus only the genetic information for half of the antigen-binding site. However, the genes for both antibody chains can be recovered separately by plating on the selective medium, suggesting a means by which mutually complementary pairs of antigen binding heavy and light chain combinations could be selected from random combinatorial libraries. For example, a light chain repertoire on fd phage could be used to infect cells harbouring a library of soluble heavy chains on the phagemid. The affinity purified phagemid library could then be used to infect E.coli, rescued with the affinity purified phage library, and the new combinatorial library subjected to a further round of selection. Thus, antibody heavy and light chain genes are reshuffled after each round of purification. Finally, after several rounds, infected bacteria could be plated and screened individually for antigen-binding phage. Such 'dual' combinatorial libraries are potentially more diverse than those encoded on a single vector. By combining separate libraries of 10^7 light chain phage(mid)s, the diversity of displayed Fab fragments (potentially 10^{14}) is limited only by the number of bacteria (10^{12} per litre). More simply, the use of two vectors should also facilitate the construction of 'hierarchical' libraries, in which a fixed heavy or light chain is paired with a library or partners (example 18), offering a means of 'fine-tuning' antibody affinity and specificity.

Example 23

Induction of Soluble scFv and Fab Fragments using Phagemid pHEN1

[0364] Further study of antibodies which have been expressed on the surface of phage would be greatly facilitated if it is simple to switch to expression in solution.

[0365] E.coli HB2151 was infected with pHEN phagemid (pHEN1-I or II), and plated on YTE, 100 $\mu\text{g}/\text{ml}$ ampicillin plates. Colonies were shaken at 37°C in 2xTY medium, 100 $\mu\text{g}/\text{ml}$ ampicillin, 1% glucose to $\text{OD}_{550}=0.5$ to 1.0. Cells were pelleted, washed once in 2xTY medium, resuspended in medium with 100 $\mu\text{g}/\text{ml}$ ampicillin, 1 mM isopropyl β -D-thiogalactoside (IPTG), and grown for a further 16 hours. Cells were pelleted and the supernatant, containing the secreted chains, used directly in ELISA.

[0366] The phagemid pHEN1 has the advantage over phage fd-CAT2, in that antibody can be produced either for phage display (by growth in supE strains of E.coli) or as a tagged soluble fragment (by growth in non-suppressor strains), as a peptide tag (example 20) and amber codon were introduced between the antibody and g3p. Secretion of soluble Fab fragments from pHEN1-II or scFv fragments from pHEN1-I was demonstrated after growth in E.coli HB2151 and induction with IPTG using Western blots (Figure 26). For detection of secreted proteins, 10 μl supernatant of induced cultures were subjected to SDS-PAGE and proteins transferred by electroblotting to Immobilon-P (Millipore). Soluble heavy and light chain were detected with goat polyclonal anti-human Fab antiserum (Sigma) and peroxidase conjugated rabbit anti-goat immunoglobulin (Sigma), each at a dilution of 1:1000. The tagged VK domain was detected with 9E10 antibody (1:1000) and peroxidase conjugated goat anti-mouse immunoglobulin (Fc specific) (1:1000) (Sigma) or with a peroxidase labelled anti-human CK antiserum (Dako). 3,3'-diaminobenzidine (DAB;Sigma) was used as peroxidase substrate (Harlow E., et al. 1988 Supr). With the scFv, the fragments were detected using the 9E10 anti-myc tag antibody (data not shown). With the Fab, only the light chain was detected by 9E10 (or anti-human CK) antibody, as expected, while the anti-human Fab antiserum detected both heavy and light chains. Binding of the soluble scFv and Fab fragments to pHx-BSA (but not to BSA) was also demonstrated by ELISA (Table 4B). Thus scFv and Fab fragments can be displayed on phage or secreted as soluble fragments from the same phagemid vector.

Example 24

Increased Sensitivity in ELISA assay of Lysozyme using FDTscFvD1.3 as Primary Antibody Compared to Soluble scFvD1.3

[0367] In principle the use of phage antibodies should allow more sensitive immunoassays to be performed than with soluble antibodies. Phage antibodies combine the ability to bind a specific antigen with the potential for amplification through the presence of multiple (ca.2800) copies of the major coat protein (g8p) on each virion. This would allow the

attachment of several antibody molecules directed against M13 to each virion followed by the attachment of several molecules of peroxidase-conjugated anti-species antibody (anti-sheep) IgG in the case below). Thus for every phage antibody bound to antigen there is the potential for attaching several peroxidase molecules whereas when a soluble antibody is used as the primary antibody this amplification will not occur.

[0368] ELISA plates were coated overnight at room temperature using 200 μ l of 10 fold dilutions of hen egg lysozyme (1000, 100, 10, 1, 0.1 and 0.01 μ g/ml) in 50mM NaHCO₃, pH9.6. ELISA was performed as described in example 4 except that (i) incubation with anti-lysozyme antibody was with either FDTscFvD1.3 (pAb; 10¹¹ phage per well; 1.6mol) or soluble affinity purified scFvD1.3 (18 μ g per well; 0.7nmol) (ii) incubation with second antibody was with 1/100 dilution of sheep anti-M13 serum for FDTscFvD1.3 samples or with or 1/100 dilution of rabbit anti-scFvD1.3 serum (from S. Ward)- for soluble scFvD1.3 samples (iii) peroxidase-conjugated rabbit anti-goat immunoglobulin (Sigma; 1/5000) was used for FDTscFvD1.3 samples and peroxidase-conjugated goat anti-rabbit immunoglobulin (Sigma; 1/5000) was used for soluble scFvD1.3 samples. Absorbance at 405nm was measured after 15h. The results are shown in Figures 30 and 31. In these figures lysozyme concentrations for coating are shown on a log scale of dilutions relative to 1 μ g/ml. (i.e. log = -3 = 1mg/ml ; log = 2 = 0.01 μ g/ml)

[0369] Higher signals were obtained with FDTscFvD1.3 at all concentrations of lysozyme (Fig.28) but the difference was very marked at the greatest dilutions, where antigen quantities are most limiting (Figs. 27 and 28). This suggests that phage antibodies may be particularly valuable for sandwich type assays where the capture of small amounts of antigen by the primary antibody will generate an amplified signal when phage antibodies directed against a different epitope are used as the second antigen binding antibody.

Example 25

Direct Rescue and Expression of Mouse Monoclonal Antibodies as Single Chain Fv Fragments on the Surface of Bacteriophage fd.

[0370] The principle is very similar to that described in example 12. It consists of the PCR assembly of single chain antibodies from cDNA prepared from mouse monoclonals. As an example, the rescue and expression of two such antibodies from monoclonals expressing antibodies against the steroid hormone oestriol is described.

A. RNA Preparation

[0371] RNA can be prepared using many procedures well known to those skilled in the art. In this example, the use of Triton X-100 lysis, phenol/SDS RNase inactivation gave excellent results.

1. The mouse monoclonal cells that were used here had been harvested by centrifugation and resuspended in serum free medium. They were then centrifuged and resuspended in saline and after a final centrifugation step, resuspended in sterile water at 1 x 10⁷ cells per ml. (Normally cells would be washed in PBS buffer and finally resuspended in PBS buffer, but these particular cells were supplied to us as described frozen in water.).

2. To 750 μ l of cells was added 250 μ l of ice cold 4X lysis buffer (40mM Tris HCl pH 7.4/4mM MgCl₂/600mM NaCl/40mM VRC (Veronyl ribosyl complex)/2% Triton X-100). The suspension was mixed well and left on ice for 5 minutes.

3. Centrifugation was carried out at 4°C in a microfuge at 13000 rpm for 5 min.

The supernatant is then phenol extracted three times, phenol chloroform extracted three times and finally, ethanol precipitated as described in the materials and methods. The precipitate was resuspended in 50 μ l water.

4. The optical density of the RNA at 260nm with a 2.5 μ l sample in 1ml water was measured. The RNA was checked by electrophoresis of a 2 μ g sample on a 1% agarose gel. RNA in the range of 32 μ g to 42 μ g was obtained by this method.

B. cDNA Preparation

[0372] The method used is the same as that described in example 12. Two cDNA preparations were made. These were from RNA extracted from the monoclonals known as cell lines 013 and 014 which both express antibodies against the steroid hormone, oestriol.

C. Primary PCRs

[0373] The method used is essentially the same as that described in example 14. The VH region was amplified with the primers VH1BACK and VH1FOR-2. For the V κ region, four separate reactions were carried out using the

primer VK2BACK and with MJK1FONX, MJK2FONX, MJK4FONX or MJK5FONX. Samples (5ul) were checked on a 1.5% agarose gel. From this it was observed that for cDNA prepared from the two oestriol monoclonals the primers VK2BACK and MJK1FONX gave the best amplification of the V κ region. The VH bands and the V κ bands amplified with VK2BACK/MJK1FONX were purified on 2% low melting point agarose gels for each monoclonals. The DNA bands were excised from the gel and purified using a dedicated GeneClean kit as described in example 12.

D. Preparation of linker

[0374] The method used is essentially the same as that described in example 12. In this case, the amplified linker DNA was purified on a 2% agarose gel and recovered from the gel with a dedicated "Mermaid" kit (BIO 101, GeneClean, La Jolla, San Diego, California, USA) using the manufacturers instructions.

E. Assembly PCRs

[0375] The method used is essentially the same as that described in example 12. In this case, the assembled PCR product was purified on a 2% agarose gel and recovered from the gel with a dedicated "Mermaid" kit.

F. Adding restriction sites and work-up

[0376] The assembled product was "tagged" with Apa LI and Not I restriction sites. The DNA was then digested with Apa LI and Not I to give the appropriate sticky ends for cloning and then purified on a 2% low melting point agarose gel and extracted using a GeneClean kit. The method used is the same as that described in example 12.

G. Cloning into Vector fd-CAT2

[0377] A total of 15ug of CsCl purified fd-CAT2 DNA was digested with 100 units of the restriction enzyme Not I (New England Biolabs) in a total volume of 200ul 1X NEB Not I buffer with 1X NEB acetylated BSA for a total of 3 hours at 37°C. The vector DNA was treated twice with 15ul StrataClean (a commercially available resin for the removal of protein), following the manufacturers instructions (Stratagene, 11099 North Torrey Pines Road, La Jolla, California, USA). The DNA was then ethanol precipitated and redissolved in TE buffer (Sambrook et al., 1989 supra). The DNA was then digested with 100 units of the restriction enzyme Apa LI (New England Biolabs) in a total volume of 200ul 1X NEB Buffer 4 overnight at 37°C. The vector was then purified with a Chroma Spin 1000 column following the manufacturers instructions (Clontech Laboratories Inc, 4030 Fabian way, Palo Alto, California, USA). This step removes the Apa LI/Not I fragment to give cut vector DNA for maximum ligation efficiency.

[0378] Ligation reactions were carried out with 2.5-long of the DNA insert and 10ng of vector in a total volume of 10ul of 1X NEB ligase buffer with 1ul of NEB ligase (New England Biolabs) at 16°C overnight (approx 16 hours).

H. Transformation and growth

[0379] E.coli strain TG1 was made competent and transformed with the fdCAT2 recombinant DNA as described by Sambrook et al, 1989 Supra. The cells were plated out on LBtet plates (10g tryptone, 5g yeast extract, 10g NaCl, 15g bacto-agar per litre with 15ug/ul of tetracycline added just before pouring the plates) and grown overnight.

[0380] Single well isolated colonies were then inoculated into 10 ml of LBtet broth (LB medium with 15ug/ul of tetracycline) in 50 ml tubes. After overnight growth at 35°C/350rpm in a bench top centrifuge. The supernatants were transferred to 15 ml centrifuge tubes and 2ml 20% PEG 8000/2.5M NaCl added to each. After incubating at room temperature for 20-30 minutes, the recombinant phage was pelleted by centrifugation at 9000rpm in a Sorval SM24 rotor for 30 minutes. The PEG supernatant was discarded. Any remaining PEG was removed with a pasteur pipette after a brief (2 minutes) centrifugation step. This last step was repeated to make sure that no PEG remained. The phage pellet was then resuspended in 500ul PBS buffer. This was transferred to a microcentrifuge tube and spun at 13000 rpm to remove any remaining cells. The phage supernatant was transferred to a fresh tube.

I. Assay for antibody expression

[0381] Bacteriophage fd recombinants were screened for the expression of antibody against oestriol by ELISA. This method is described in example 6. In this case the following alterations are relevant.

1. Microtitre plates were coated overnight with 40ug/ml oestriol-6 carboxymethyloxime-BSA (Steraloids, 31 Radcliffe Road, Croydon, CRO 5QJ, England).

2. 1st antibody was the putative phage anti oestriol antibody. 50ul of phage in a final volume of 200ul of sterile PBS combining 0.25% gelatin was added to each well.
3. 2nd antibody was sheep anti M13 at 1:1000 dilution.
4. 3rd antibody was peroxidase conjugated rabbit anti goat immunoglobulin.

[0382] Recombinants expressing functional antibody were detected by incubation with the chromogenic substrate 2'2' zindois (3-ethyl benzthiazoline sulphonc acid). The results are shown in figures 29 and 30.

Example 26

PCR Assembly of DNA encoding Fab Fragments of an Antibody directed against Oxazolone

[0383] Example 21 showed that genes encoding Fab fragments could be subcloned into vectors fdCAT2 and pHEN1 and the protein domains displayed on the surface of phage with retention of binding function. This example shows that the VHCH and VKCK domains can be amplified separately and then joined by a linker allowing the expression of the light chain as a geneIII protein fusion and the VHCH fragment as a soluble molecule. A functional Fab fragment is then displayed on phage by association of these domains. The assembly process, described in this example, is required for display of a library of Fab fragments derived from the immune repertoire if both heavy and light chain domains are to be encoded within a single vector.

[0384] The VHCH1 and VKCK domains of a construct (example 21; construct II in pUC19) derived from antibody. NQ10 12.5 directed against 2-phenyl-5-oxazolone were amplified using PCR. For cloning into the vector fdCAT2 the oligonucleotides VH1BACKAPA (example 21) and HulgI-4 CH1FOR (example 30) were used to amplify the VHCH1 domains. For cloning into pHEN1 VH1BACKSFH5 (example 21) replaced VH1BACKAPA for this amplification. For cloning into both vectors the VKCK domains were amplified using VK2BACK (example 21) and CKNOTFOR (example 30). A linker oligonucleotide fragment containing the bacteriophage fd gene 8 terminator and the fd gene 3 promoter was prepared by amplifying the region containing them from the vector fdCAT2 by PCR using the oligonucleotides.

VK-TERM-FOR

5' TGG AGA CTG GGT GAG CTC AAT GTC GGA GTG AGA ATA GAA

AGG 3' (overlapping with VK2BACK [example 12])
and

**5' AAG CCC AGC AAC ACC AAG GTG GAC AAG AAA GTT GAG CCC AAA
TCT AGC TGA TAA AÇC GAT ACA ATT AAA GGC 3' (overlapping
with HuIgG1-4 CH1-FOR)**

Assembly of the Fab fragment from the amplified VHCH1 and VKCK domains and the linker prepared as above was as described in example 12E except that the primers VH1BACKAPA (when cloning into fdCAT2) or VH1BACKSFH5 (when cloning into pHEN1) and CKNOTFOR were used for the final reamplification, thereby introducing restriction sites for cloning into fdCAT2 (Apa1I-NotI) or pHEN1 (SfiI-NotI) the assembled Fab fragment is shown in figure 31. No assembled product was seen in the absence of linker. An assembled scFv prepared according to example 12 is shown for comparison.

[0385] Phage antibodies were prepared as in example 21 and ELISA was performed with oxazolone as antigen according to example 6. Results were as expected for Fab fragments cloned in both fdCAT2 and pHEN1 samples, phage particles bound to oxazolone as detected by a positive ELISA signal.

Example 27

Construction of a Gene III Deficient Helper Phage

[0386] To fully realise the potential of the phagemid cloning system, a helper phage lacking gene III is desirable. Rescue of gene III fusions with such a helper phage would result in all the progeny phagemids having a gene III fusion on their capsid, since there would be no competition with the wild type molecule.

[0387] Control over the number of fusion molecules contained on each phage will provide particularly useful. For example, a gene III deficient helper phage can be used to rescue low affinity antibodies from a naive repertoire, in which high avidity will be necessary to isolate those phage bearing the correct antibody specificity. The unmutated

helper phage can then be used when higher affinity versions are constructed, thereby reducing the avidity component, and permitting selection purely on the basis of affinity. This will prove a surprisingly successful strategy for isolation and affinity maturation of antibodies from naive libraries.

5 **[0388]** The strategy chosen to construct the helper phage was to partially delete gene III of M13K07 using exonuclease Bal 31. However, phage lacking gene III protein are non-infective so an E.coli strain expressing gene III was constructed. Wild type M13 gene III was PCR-amplified with primers gIIIFUFO and gIIIFUBA, exactly as described in example 20. The PCR product was digested with Eco RI and Hind III and inserted into Eco RI and Hind III-cut pUC19 (not a phagemid as it lacks the filamentous phage origin of SS DNA replication) under control of the lac promoter. The plasmid was transformed into E.coli TG1, and the resulting strain called TG1/pUC19gIII. This strain provides gIII protein

10 in trans to the helper phage.
[0389] There is a single unique Bam HI site in M13K07, which is approximately in the centre of gIII. Doublestranded M13K07 DNA was prepared by alkaline lysis and caesium chloride centrifugation (Sambrook et al, et supra. 1989); twenty µg of DNA was cut with Bam H1, phenol extracted and ethanol precipitated then resuspended in 50µl of Bal 31 buffer (600mM NaCl, 20mM Tris-HCl pH 8.0, 12 mM CaCl₂, 12mM MgCl₂ and 1mM EDTA) and digested for 4 minutes

15 with 1 unit of Bal 31 (New England BioLabs). This treatment removed approximately 1Kb of DNA. EGTA was added to 20mM and the reaction phenol extracted and ethanol precipitated prior to purification of the truncated genome on an agarose gel. The DNA was repaired with klenow enzyme and self-ligated with T4 DNA ligase (New England BioLabs).
[0390] Aliquots of the ligation reaction were transformed into competent TG1/pUC19gIII and plated on SOB medium containing ampicillin at 100µg/ml and kanamycin at 50µg/ml. Colonies were screened for the presence of a deletion

20 by PCR with primers gIIIFUBA and KSJ12 (CGGAATACCCAAAAGAACTGG).
[0391] KSJ 12 anneals to gene VI which is immediately downstream of gIII in the phage genome, so distinguishing gIII on the helper phage from that resident on the plasmid. Three clones gave truncated PCR products corresponding to deletions of ca. 200, 400 and 800bp. These clones were called M13K07 gIII Δ Nos 1, 2 and 3 respectively. No clones were isolated from the earlier Bal 31 time points, suggesting that these are in some way lethal to the host cell. Several

25 clones were isolated from later time points, but none of these gave a PCR product, indicating that the deletion reaction had gone too far.
[0392] M13K07 gIII Δ Nos 1, 2 and 3 were cultured and the resulting helper phage tested for their ability to rescue an antibody gIII fusion (scFv D1.3) by ELISA, exactly as described in example 14. As shown in figure 32, only one clone, M13K07 gIIIΔ No3 was found to rescue the antibody well; in fact the signal using this helper was greater than that observed with the parent M13 K07. M13K07 gIIIΔ No3 rescued phagemids should have a much higher density of antibody fusions on their surfaces. That this was indeed the case was demonstrated when the phage used in this

30 ELISA were analysed by Western blotting with anti gIII protein antiserum (fig.33). This analysis enables estimation of the amount of gIII fusion protein versus free gIII protein present on the phage(mid) particles.
[0393] Only a minute fraction of the gIII protein on the M13K07-rescued material is present as an intact fusion (fig. 33). The fusion protein band is induced by IPTG, so is indisputably that synthesised by the phagemid. As expected, even when the lac promoter driving gIII fusion protein synthesis is fully induced (100µM IPTG), wild type gIII protein, at a lower copy number and driven from a far weaker promoter, predominates. This is in contrast to the pattern generated by the same clone rescued with M13K07 gIIIΔNo3, and the pattern generated by fd CAT2-scFv D1.3. In both of these latter cases, there is no competition with wild-type gIII and the fusion protein band is correspondingly stronger.

40 **[0394]** It is worthy of note that construction of M13K07 gIII Δ No3 was immensely inefficient: one clone from 20µg of starting DNA. Moreover, the yield of gIII helper phage from overnight cultures is extremely low ca. 10⁶ cfu/ml compared with ca. 10¹¹ cfu/ml for the parental phage. Despite this, M13K07 gIII No3 rescues the phagemid as well as the parental phage, as judged by the number of phagemid particles produced after overnight growth. This indicates that trans replication and packaging functions of the helper are intact and suggest that its own replication is defective. Hence it may be that inactivation of gIII is normally toxic to the host cell, and that M13K07 gIIIΔNo3 was isolated because of a compensating mutation affecting, for example, replication. Phage fd-tet is unusual in that it tolerates mutations in structural genes that are normally lethal to the host cell, since it has a replication defect that slows down accumulation of toxic phage products; M13K07 gIIIΔ No3 may also have such a defect.

45 **[0395]** M13K07g IIIΔ No 3 has been deposited at the National Collection of Type Cultures, 61 Colindale Avenue, London, NW9 6HT, UK (Accession No. NCTC 12478). On 28 June 1991, in accordance with the regulations of the Budapest Treaty. It contains a deletion of the M13 genome from bases 1979 to 2768 inclusive (see Van Wezenbeek, P.G.M.F. et al., Gene 112:129-148, 1980 for the DNA sequence of the M13 genome).

Example 28Selection of bacteriophage expressing scFv fragments directed against lysozyme from mixtures according to affinity using a panning procedure

[0396] For isolation of an antibody with a desired high affinity, it is necessary to be able to select an antibody with only a few fold higher affinity than the remainder of the population. This will be particularly important when an antibody with insufficient affinity has been isolated, for example, from a repertoire derived from an immunised animal, and random mutagenesis is used to prepare derivatives with potentially increased affinity. In this example, mixtures of phage expressing antibodies of different affinities directed against hen egg lysozyme were subjected to a panning procedure. It is demonstrated that phage antibodies give the ability to select for an antibody with a K_d of 2nM against one with a K_d of 13nM.

[0397] The oligonucleotides used in this example are shown in the list below:

OLIGONUCLEOTIDES

VHBHD13APA : 5' - CAC AGT GCA CAG GTC CAA CTG CAG GAG AGC
GGT

VHFHD13 : 5' - CGG TGA CGA GGC TGC CTT GAC CCC

HD13BLIN : 5' - GGG GTC AGG GCA GCC TCG TCA CCG

HD13FLIN3 : 5' - TGG GCT CTG GGT CAT CTG GAT GTC CGA T

VKBHD13 : 5' - GAC ATC CAG ATG ACC CAG AGC CCA

VKFHD13NOT : 5' - GAG TCA TTC TGC GGC CGC ACG TTT GAT TTC
CAC CTT GGT CCC

MURD13SEQ : 5' - GAG GAG ATT TTC CCT GT

HUMD13SEQ : 5' - TTG GAG CCT TAC CTG GC

FDPCRFOR : 5' - TAG CCC CCT TAT TAG CGT TTG CCA

FDPCRBK : 5' - GCG ATG GGT GTT GTC ATT GTC GGC

Phage displaying scFv fragments directed against lysozyme were derived from cloned Fab fragments in plasmids.

[0398] Heavy and light chain variable regions were amplified by the polymerase chain reaction (PCR) from plasmids containing humanized VH-CH1 or VK-CK inserts suitable for production of Fab fragments (gift of J. Foote). The dissociation constant, K_d for different combinations of the two plasmids combined as Fabs, are shown below:

Heavy Chain Plasmid	Light Chain Plasmid	K_d
HuH-1	HuK-3	52 nM
HuH-1	HuK-4	180 nM
HuH-2	HuK-3	13 nM
HuH-2	HuK-4	(not determined)

Primary PCR

[0399] The primary PCR of the variable regions was performed by combining the following:

36.5 μ l Water
5 μ l PCR buffer (10x)
2 μ l dNTP (5mM)
2.5 μ l Back oligo (10 pmoles/ μ l) (VHBHD13APA or VKBHD13)
2.5 μ l Forward oligo (10 pmoles/ μ l) (VHFHD13 or VKFHD13NOT)

[0400] The reaction is decontaminated by UV irradiation to destroy foreign DNA for 5 minutes, and 1 μ l of plasmid DNA added (0.1 μ g/ μ l). The PCR mixture was covered with 2 drops of paraffin oil, and placed on the PCR block at

94°C for 5 minutes before the addition of 0.5 µl of Taq DNA polymerase under the paraffin. The cycling conditions used were 94°C 1 min, 40°C 1 min, 72°C 1.5 min 17 cycles.

[0401] The linker (Gly₄-Ser)₃, was amplified from the anti-phOx (2-phenyloxazol-5-one) clone fd-CAT2-scFv NQ11, using the oligos HD13BLIN and HD13FLIN3, with 0.1 µg of plasmid DNA. The PCR cycling used was 94°C 1 min, 25°C 1.5 min, for 17 cycles.

[0402] Amplified DNA was purified by running the samples on a 2% low melting point agarose gel at 90 mA, excising the appropriate bands and extracting the DNA using the GeneClean II Kit (BIO 101 Inc.) for the VH and VK, or by using Spin-X filter units (Costar) for the linker. A final volume of 10 µl was used to resuspend the extracted DNA.

PCR Assembly

[0403] Assembly of the four single chain Fv Humanized D1.3 (scFv HuD1.3) constructs was by the process of 'assembly by overlap extension' example 12. The following were combined:

34.5 µl Water
5 µl PCR Buffer (10x)
2 µl dNTP (5 mM)
2.5 µl Back oligo (10 pmoles/µl) (VHBHD13APA)
2.5 µl Forward oligo (10 pmoles/µl) (VKFHD13NOT)

[0404] Once again, the reaction is decontaminate by UV treatment for 5 minutes before the addition of 1 µl of the primary PCR products; VH-1 or VH-2, VK-3 or VK-4, plus the linker DNA. The reaction was covered with 2 drops of paraffin, and heated at 94°C for 5 minutes before the addition of 0.5 µl of Taq Polymerase. The PCR cycling conditions used were 94°C 1 min, 60°C 1.5 min, 72°C 2.5 min for 20 cycles.

[0405] The aqueous layer under the paraffin was extracted once with phenol, once with phenol: chloroform, once with ether, ethanol precipitated, and resuspended in 36 µl of water. To this was added, 5 µl of 10x Buffer for NotI, 5 µl 1 mg/ml BSA, and 4 µl (40 U) of NotI (New England Biolabs). The restriction was incubated at 37°C overnight.

[0406] The DNA was ethanol precipitated and resuspended in 36 µl of water, and 5 µl 10x NEB Buffer 4, 5 µl 1 mg/ml BSA, and 2 µl (40 U) of ApaLI (New England Biolabs). This was incubated at 37°C for 5 hours; a further 2 µl of ApaLI was added and the reaction incubated at 37°C overnight.

[0407] The cut DNA was extracted by gel purification on a 1.3% low melting point agarose gel followed by treatment with GeneClean, to yield the insert DNA for cloning.

[0408] Vector fd CAT2 (prepared and digested with ApaLI and NotI as in example 16) and the scFv DNA were ligated as in example 16.

Analysis Of Clones

[0409] Colonies from the ligations were first screened for inserts by PCR screening. The PCR mixture was prepared in bulk by combining 14.8 µL 1x PCR Buffer, 1 µl dNTP (5 mM), 1 µl Back oligo (FDPCRBK), 1 µl Forward oligo (FDPCRFOR), and 0.2 µl Taq polymerase per colony screened. 20 µl of this PCR mixture was aliquoted into a 96 well Techn plate. The top of a colony was touched with a toothpick and twirled quickly into the PCR mixture and the colony rescued by placing the toothpick in a Cellwell plate (Nunc) containing 250 µl of 2x TY medium. The PCR mixture is covered with 1 drop of paraffin and the plate placed on the block at 94°C for 10 minutes before cycling at 94°C 1 minute, 60°C 1 minute, 72°C 2.5 minutes.

[0410] The clones thus derived were named as below. The affinity of scFv fragments derived from the Fab fragments was not determined but previous results suggests that these are closely related although not necessarily identical (R. E. Bird & B.W. Walker TIBTECH 9 132-137, 1991).

Construct Name	Composition	Affinity of Fab (Kd)
TPB1	VH-HuH2-(Gly ₄ -Ser) ₃ -VK-HuK3	13 nM
TPB2	VH-HuH1-(Gly ₄ -Ser) ₃ -VK-HuK4	180 nM
TPB3	VH-HuH2-(Gly ₄ -Ser) ₃ -VK-HuK4	(Unknown)
TPB4	VH-HuH1-(Gly ₄ -Ser) ₃ -VK-HuK3	52 nM

[0411] Preparation of phage and ELISA was as described in example 6. The clones generated in fd CAT2 were shown to bind lysozyme as expected.

Affinity selection

Selection of Highest Affinity Binding Phage

5 **[0412]** Mixing experiments were performed in which fd-CAT2 scFvD1.3 phage (example 15) were mixed with either fd-CAT2 TPB1, fd-CAT2 TPB2, or fd-CAT2 TKPB4, and used in one round of panning.

[0413] The general method used for affinity selection by panning is that detailed below. Any deviation from this protocol is described at the relevant point. Panning plates were placed on a rocking platform between manipulations.

10 **[0414]** Falcon 35 mm Tissue Culture dishes were coated overnight with 1 ml of Lysozyme (various concentrations) dissolved in 50 mM Sodium Hydrogen Carbonate, pH 9.6, and blocked with 2 ml 2% MPBS at room temperature for 2 hours. Phage were prepared in 1 ml 2% MPBS and rocked at room temperature for 2 hours. Plates were washed for 5 minutes with 2 ml of the following solutions; 5 times with PBS, PBS-Tween, 50 mM Tris-HCl, pH 7.5; 500 mM Sodium Chloride, 50 mM Tris-HCl, pH 8.5; 500 mM Sodium Chloride, 50 mM Tris-HCl, pH 9.5; 500 mM Sodium Chloride, 50 mM Sodium Hydrogen Carbonate, pH 9.6; 500 mM Sodium Chloride. Phage were then eluted by adding 1 ml 100 mM Triethylamine and rocking for 5 minutes before removing the eluate which was neutralised with 100 µl 1.0 M Tris-HCl, pH 7.4.

[0415] Plates were coated overnight with Lysozyme at the concentration listed below.

[0416] Colonies from the single round of panning were probed with either MURDSEQ (for fdCAT2 scFvD1.3) or HUMD13SEQ (for fdCAT2 TPB constructs).

20 **[0417]** Circles of nitrocellulose (Schleicher & Schuell, BA 85, 0.45 µm) were labelled in pencil and lowered gently onto the colonies derived from the panning experiments and left for one minute. The filters were then pulled off quickly from one edge and placed colony side up on a piece of 3MM paper (Whatman) soaked in Denaturing solution (500 mM Sodium Hydroxide; 1.5 M Sodium Chloride) for 5 minutes. They were then transferred to 3MM soaked in Neutralizing Solution (3.0 M Sodium Chloride; 500 mM Tris-HCl, pH 7.5) for 1 minute, and then to 3MM soaked in 5x SSC; 250 mM Ammonium Acetate for 1 minute. The filters were then air dried before baking in an 80°C vacuum oven for 30 minutes.

[0418] The oligonucleotide probe was prepared by combining the following:

30 2 µl oligonucleotide (1 pmoles/µl)
2 µl γ-32P ATP (3000 Ci/mmol) (Amersham International plc)
2 µl 10 x Kinase buffer (0.5 M Tris-HCl, pH 7.5; 100 mM Magnesium Chloride; 10 mM DTT)
12 µl Water
2 µl Polynucleotide Kinase (20 Units)

35 This was incubated at 37°C for 1 hour.

[0419] Hybridization was performed in the Techne HB-1 Hybridiser. The baked filters were pre-hybridized at 37°C in 40 ml of Hybridization Buffer (10 ml 100 mM Sodium pyrophosphate; 180 ml 5.0 M Sodium chloride; 20 ml 50x Denharts Solution; 90 ml 1.0 M Tris-HCl, pH 7.5; 24 ml 250 mM EDTA; 50 ml 10% NP40; made to 1 litre with water; 60.3 mg rATP; 200 mg yeast RNA (Sigma)), for 15 minutes before the addition of the 20 µl of the kinased oligo. The filters were incubated at 37°C for at least one hour, and then washed 3 times with 50 ml of 6x SSC at 37°C for 10 minutes (low stringency wash). Filters were air dried, covered with Saran wrap and exposed overnight with Kodak X-AR film.

Selection of fd-CAT2 scFv D1.3 from fd-CAT2 TPB4

45 **[0420]** Figure 34, summarizes the results from panning experiments using a mixture of the high affinity fd-CAT2 scFv D1.3 phage (Kd-2 nM) and the fd-CAT2 TPB4 construct (Kd-52 nM).

[0421] At a coating concentration of 3000 µg/ml Lysozyme, little or no enrichment could be obtained. It was however, possible to get enrichment for the scFv D1.3 phage when a lower concentration of Lysozyme was used for coating the plates. The best enrichment value obtained was from 1.5% fd-CAT2 scFv D1.3 in the starting mixture, to 33% fd-CAT2 scFv D1.3 in the eluted fraction, on a plate coated overnight with 30 µg/ml Lysozyme.

Selection of fd-CAT2 scFv D1.3 from fd-CAT2 TPB1

55 **[0422]** Enrichment for the high affinity scFv D1.3 phage over the fd-CAT2 TPB1 phage (Kd-13) nM, could only be shown from experiments where the plates had been coated overnight with low concentrations of Lysozyme, as shown in Figure 35.

[0423] In summary, single chain Fv versions of a series of humanized D1.3 antibodies have been constructed in phage fd-CAT2. By affinity selection of fd-CAT2 phage mixtures, by panning in small petri dishes, it was shown that

the high affinity scFv D1.3 phage, could be preferentially selected for against a background of lower affinity scFv HuD1.3 phage.

Example 29 Generation and Selection of Mutants of an Anti-4-hydroxy-3-nitrophenylacetic acid (NP) Antibody expressed on Phage using Mutator strains

[0424] It will sometimes be desirable to increase the diversity of a pool of genes cloned in phage, for example a pool of antibody genes, or to produce a large number of variants of a single cloned gene. There are many suitable in vitro mutagenesis methods. However, an attractive method, particularly for making a more diverse population of a library of antibody genes, is to use mutator strains. This has the advantage of generating very large numbers of mutants, essentially limited only by the number of phage that can be handled. The phage display system allows full advantage to be taken of this number to isolate improved or altered clones.

[0425] Nucleotide sequences encoding an antibody scFv fragment directed against 4-hydroxy-3-nitrophenylacetic acid (NP), scFvB18, derived as in example 12, from a monoclonal antibody against NP were cloned into fdCAT2 using ApaI and NotI restriction sites as in example 11 to create fdCAT2scFvB18 or into fdDOGkan (fdCAT2 with its tetracycline resistance gene removed and replaced by a kanamycin resistance gene) using PstI and NotI restriction sites to create fdDOGkanscFvB18 or into the phagemid vector pHEN1 using the restriction sites SfiI and NotI as a fusion protein with gene III to create pHEN1scFvB18.

[0426] The following mutator strains (R. M. Schaaper & R.L. Dunn J. Mol. Biol. 262 1627-16270, 1987; R. M. Schaaper Proc. Natl. Acad. Sci. U.S.A. 85 8126-8130 1988) were used:

NR9232: ara, thi, mutD5-zaf13::Tn10, prolac, F'prolac
 NR9670: ara, thi, azi, mutT1, leu::Tn10, prolac
 NR9292: ara, thi, mutH101, prolac, F'prolac
 NR9084: ara, thi, mutT1, azi, prolac, F'prolac⁺ΔM15 M15
 NR9046: ara, thi, supE, rif, nalA, metB, argE(am), prolac, F'prolac
 were kind gifts of Dr. R. M. Schaaper (Department of
 Health & Human Services, NIH, PO Box 12233, Research Triangle Park, N.C. 27709)
 NR9046mutD5: NR9046 mutD5::Tn10
 NR9046mutT1: NR9046 mutT1::Tn10

were constructed by P1 transduction according to standard procedures. Mutator strains were transfected with fdCAT2scFvB18 or fdDOGkanscFvB18 and transfectants selected for antibiotic resistance. Transfectants were grown for 24h at 37°C before mutant phage was harvested by PEG precipitation. The mutant phage were selected on a 1ml NIP (4-hydroxy-3-iodo-5-nitrophenylacetic acid)-BSA-Sepharose affinity column (prepared according to the manufacturers instructions) prewashed with 200ml of PBS and blocked by 20ml MPBS. Phage were loaded on the column in 10ml MPBS and unbound material reapplied to ensure complete binding. The column was subsequently washed with 10ml of MPBS and 500ml of PBS. Phage bound to the affinity matrix was eluted with 5 column volumes of 0.33 mM NIP-Cap (example 38).

[0427] Phage eluate was incubated for 30min to 1h with log phase (2×10^8 cells/ml) E.coli mutator strains without antibiotic selection. The infected cells were then diluted 1:100 in 2xTY and grown for 24h with antibiotic selection (15μg/ml tetracycline or 30μg/ml kanamycin for fdCAT2scFvB18 or fdDOGkanscFvB18 respectively). Phage from this culture was used for another round of affinity selection and mutation.

[0428] Binding of phage antibodies was assayed by ELISA as in example 9 except that ELISA plates were coated with NIP-BSA (4-hydroxy-3-iodo-5-nitrophenylacetyl-BSA; 0.4 mg/ml). Culture supernatants were prepared following growth in Cellwells as described in example 17 and 20μl of culture supernatant was added to each well diluted to 200μl with MPBS.

[0429] Phage samples giving signals in ELISA of more than twice the background were tested ELISA as above for non-specific binding against lysozyme, BSA or Ox-BSA (example 9). Specificity for NIP was further confirmed by an ELISA in which serial dilutions of NIP-CAP were added together with phage antibodies. Addition of increasing concentrations of NIP-CAP reduced the ELISA signal to the background level.

[0430] Phage giving positive signals in ELISA were sequenced and 2 different mutants were subcloned into pHEN1 phagemid and transformed into HB2151 for soluble expression and TG1 for phage display (example 23).

[0431] For expression of soluble scFv fragments, transformants in E.coli HB2151 were grown at 37°C in 1 litre 2xTY, 0.2% glucose, 0.1mg/ml ampicillin to an OD600 of 1 and expression of soluble scFv fragments induced by adding IPTG to 1mM. Cultures were shaken at 30°C for 16h.

[0432] Soluble scFvB18 was concentrated from crude bacterial supernatant in a FLOWGEN ultrafiltration unit to a volume of 200ml.

[0433] The concentrate was passed two times over a 2ml column of NIP-BSA-Sepharose prewashed with 200ml of PBS. The column was washed with 500ml of PBS and 200ml of 0.1M Tris pH7.5, 0.5M NaCl and phage antibodies eluted with 50mM Citrate buffer pH2.3. The eluate was immediately neutralised with 1MTris pH8. The eluate was dialysed against two changes of 1 litre PBS, 0.2mM EDTA, Precipitated protein was removed by centrifugation at 10000g and protein yield was determined by measuring the absorbance at 280nm of the supernatant.

[0434] After 4 rounds of mutation and selection, isolated clones were screened and in one or two rare examples strongly positive ELISA signals were obtained from phage antibodies derived from the mutation of each of fdCAT2scFvB18 and fdDOGKanscFvB18 in the ELISA. The ELISA conditions were such that the parent phage fdCAT2scFvB18 only generated weak signals. These phage antibodies giving strongly positive ELISA signals were enriched in further rounds by a factor of roughly 2.5 per round. Forty phage antibodies giving strongly positive signals were sequenced and they each displayed single mutations in six different positions in the scFvB18 nucleotide sequences, five of which reside in the light chain. More than 70% of the mutations occurred at positions 724 and 725 changing the first glycine in the J segment of the light chain (framework 4) to serine (in 21 cases) or aspartate (in 3 cases). The mutations found are shown in Table 5. The sequence of scFvB18 is shown in Figure 36.

[0435] The nucleotide sequences encoding the scFv fragments of a framework mutant with the above glycine to serine mutation, as well as a mutant where Tyr in the CDR3 of the light chain had been mutated to aspartate, were amplified by PCR from the phage antibody clones and subcloned into pHEN1 phagemid (essentially as in example 21). This avoids possible problems with genelll mutations caused by the mutator strains. The same pattern of ELISA signals was seen when the mutants were displayed on phage following rescue of the phagemid with helper phage (as described in example 21) as when the mutants were assayed when expressed from the phage genome as above.

[0436] The scFv fragments from scFvB18 and the scFv fragments containing the glycine to serine and tyrosine to aspartate mutations respectively were expressed in solution (following transformation into E.coli HB2151 as in example 23) at 30°C. They showed no differences in the ELISA signals between wild-type B18 and the framework mutant. The signal obtained from the phage antibody with the Tyr mutated to aspartate in CDR3 of scFvB18 was about 10x stronger. Expression yields were found to be comparable as judged by Western blotting using an antiserum raised against g3p (as described above). Affinity measurements were performed using fluorescence quenching as described in example 19. Affinity measurement of affinity purified scFv fragments however showed scFvB18, and the scFvB18 (Gly->Ser) and scFvB18(Tyr->Asp) mutants all to have a comparable affinity of 20nM for NIP-CAP.

[0437] A Western blot using an anti-genelll antibody showed the framework mutant had suffered significantly less proteolytic cleavage than scFvB18.

[0438] Hence, the use of mutator strains generates a diverse range of murants in phage antibodies when they are used as hosts for clones for gene III fusions. In this case some of the clones exhibit higher ELISA signals probably due to increased stability to proteolytic attack. The mutator strains can therefore be used to introduce diversity into a clone or population of clones. This diversity should generate clones with desirable characteristics such as a higher affinity or specificity. Such clones may then be selected following display of the proteins on phage.

Example 30 A PCR Based Technique for one step Cloning of Human V-genes as Fab Constructs

[0439] This example describes a PCR based technique to "assemble" human Fabs by splicing together the heavy and light chain DNA with a separate piece of 'linker' DNA. A mixture of universal primers is used which should make the technique applicable to all human V-genes.

[0440] The general technique for PCR assembly of human V-genes to create a Fab construct is described. The efficiency of this technique was assessed by "assembling", cloning and expressing a human anti rhesus-D (Rh-D) Fab from a IgG-K monoclonal hybridoma. We also demonstrate the potential to rescue human monoclonal antibodies from polyclonal cell populations by assembling, cloning, expressing and isolating an IgG-lambda monoclonal anti-Rh-D Fab from a polyclonal lymphoblastic cell line (LCL).

[0441] The overall strategy for the PCR assembly is shown in fig. 37 and is described in more detail below. For Fab assembly, the VH-CH1 and VK-CK or V lambda-C lambda light chains are amplified from first strand cDNA and gel purified. Heavy and light chain DNA are then combined together with linker DNA and flanking oligonucleotides in a new PCR reaction. This results in a full length Fab construct since the 5' end of the linker DNA is complementary to the 3' end of the CH1 domain and the 3' end of the linker is complementary to the 5' end of the light chain domain. The linker DNA contains terminal residues of the human CH1 domain, the bacterial leader sequence (pelB) for the light chain and the initial residues of the VK or V lambda light chain (fig.2). Finally, after gel purification, the Fab construct is reamplified with flanking oligonucleotides containing restriction sites for cloning.

Oligonucleotide primers: In order to develop the PCR cloning of human V genes it was necessary to design a new range of human specific oligonucleotide primers.

- 5 **[0442]** The PCR primers at the 5' end of the VH and VK and V lambda gene exon (BACK primers) are based on sequence data extracted from the Kabat database, (Kabat, E.A. et al, Sequences of Proteins of Immunological Interest. 4th Edition. US Department of Health and Human Services. 1987) the EMBL database, the literature (Chuchana, P., et al, Eur J. Immunol. 1990. 20:1317) and unpublished data. The sequence of the VH, VK and V lambda primers are given in table 1. In addition, extended VH primers with SfiI sites at the 5' end were also designed (Table 6) for adding a restriction site after assembly.
- 10 **[0443]** Table 6 also shows the 3' primers (FORWARD primers) designed for the PCR based cloning of human V genes. There are two sets of these depending on whether a Fab or scFv is to be produced. For Fab assembly, the forward primer was based at the 3' end of the CH1 domain, CK domain and Clambda domain. In addition, the CK and C2 FORWARD primers were also synthesized as extended versions with NotI sites at their 5' ends.
- 15 **[0444]** Primers complementary to the CH1 forward primers and the V kappa and V lambda back primers were synthesized to permit generation of linker DNA by PCR amplification of a plasmid template containing the Fab linker (Table 6). To ensure adequate amplification, the primers were extended into the actual linker sequence.

A RNA preparation

- 20 **[0445]** This is essentially the same as described in Example 12, but using material of human origin. In the results given in this example human hybridoma and human polyclonal lymphoblastic cell lines were used.

B cDNA preparation

- 25 **[0446]** Approximately 4 µg of total RNA in 20 µl water was heated at 65°C for 3 minutes, quenched on ice and added to a 30 µl reaction mixture resulting in a 50 µl reaction mixture containing 140 mM KCl, 50 mM Tris, HCl (pH 8.1 @ 42°C), 8 mM MgCl₂, 10 mM DTT, 500 µM deoxythymidine triphosphate 500 µM deoxycytosine triphosphate, 500 µM deoxyadenosine triphosphate and 500 µM deoxyguanosine triphosphate, 80 units of human placental RNase inhibitor and 10 pmol of the appropriate Forward primer (HulgG1-4CH1FOR, HulgMFOR, HuCKFOR, HuCLFOR). Two µl (50 units) of avian myeloblastosis virus (AMV) reverse transcriptase was added, the reaction incubated at 42°C for 1 hour, heated to 100°C for 3 minutes, quenched on ice and centrifuged for 5 minutes.
- 30

C Primary PCRs

- 35 **[0447]** For the primary PCR amplifications, an equimolar mixture of the appropriate family based BACK and FORWARD primers was used. (See specific examples 30a and 30b given later in this example). A 50 µl reaction mixture was prepared containing 5 µl of the supernatant from the cDNA synthesis, 20 pmol total concentration of the FORWARD primers, 250 µM dNTPs, 50 mM KCl, 100 mM Tris. HCl (pH 8.3), 1.5 mM MgCl₂, 175 µg/ml BSA and. 1 µl (5 units) *Thermus aquaticus* (Taq) DNA polymerase (Cetus, Emeryville, CA). The reaction mixture was overlaid with paraffin oil and subjected to 30 cycles of amplification using a Techne thermal cycler. The cycle was 94°C for 1 minute (denaturation), 57°C for 1 minute (annealing) and 72°C for 1 minute (extension). The product was analyzed by running 5 µl on a 2% agarose gel. The remainder was extracted twice with ether, twice with phenol/chloroform, ethanol precipitated and resuspended in 50 µl of H₂O.
- 40

D Preparation of linker

- 45 **[0448]** To make the Fab linker DNA, 13 separate PCR reactions were performed using HulgG1-4CH1FOR and each of the reverse VK or V lambda oligonucleotides. The template was approximately 1 ng of pJM-IFab D1.3 (fig.38) The PCR reaction reagents were as described above and the cycle was 94°:1 min, 45°:1min and 72°:1 min. The linkers were analyzed on a 4% agarose gel, purified on a 2% agarose gel, eluted from the gel on a Spin-X column and ethanol precipitated.
- 50

E Assembly PCRs

- 55 **[0449]** For PCR assembly of a human Fab approximately 1 µg of a primary heavy chain amplification and 1 µg of a primary light chain amplification were mixed with approximately 250 ng of the appropriate linker DNA in a PCR reaction mixture without primers and cycled 7 times (94°: 2 min, 72°:2.5 min) to join the fragments. The reaction mixture was then amplified for 25 cycles (94°:1 min, 68°-72°:1 min, 72°:2.5 min) after the addition of 20 pmol of the appropriate

flanking BACK and FORWARD primers.

F Adding Restriction Sites

- 5 **[0450]** The assembled products were gel purified and reamplified for 25 cycles (94°:1 min, 55°:1 min, 72°: 25min) with the flanking oligonucleotides containing the appended restriction sites. PCR buffers and NTPs were as described previously.

Specific examples of PCR assembly of human immunoglobulin genes

10

[0451]

- 15 a. PCR assembly of a Fab from a human hybridoma: the human monoclonal anti Rh-D cell lines Fog-1 (IgG-k) was derived from EBV transformation of the PBLs of a Rh-D negative blood donor immunized with Rh-D positive blood and has been previously described (Melamed, M.D., et al., J. Immunological Methods. 1987. 104:245) (Hughes-Jones N.C., et al., Biochem. J. 1990. 268:135) (Gorick, B.D. et al., Vox. Sang. 1988. 55:165) Total RNA was prepared from approximately 10^7 hybridoma cells. First strand cDNA synthesis was performed as described above using the primers HulgG1-4CH1FOR and HUCKFOR. Primary PCRs were performed for the VH-CH1 using a mixture of the 6 HuVHBACK primers and HulgG1-4CG1FOR and for the VK-CK using a mixture of the 6 HuVKBACK primers and HuCKFOR. A Fab construct was assembled as described above, restricted with SfiI and NotI, gel purified and ligated into pJM-I-Fab D1.3 restricted with SfiI and NotI. The ligation mixture was used to transform competent *E.coli* E.M.G. cells. Ninety-six clones were toothpicked into media in microtitre plate wells, grown to mid-log phase at 30°C and then expression of the Fab was induced by heat shocking at 42°C for 30 min followed by growing for 4 hours at 37°C. The ninety-six clones were then screened for anti-Rh-D activity as described below.
- 20 b. assembly of human Fabs from a polyclonal (LCL): A polyclonal LCL "OG" was derived from EBV transformation of approximately 10^7 peripheral blood lymphocytes (PBLs) from a Rh-D negative donor immunized with Rh-D positive red blood cells. The cells were plated at a concentration of approximately 10^5 cells per well. Positive wells were identified by screening the cells harvested and then subcloned once. Typing of the well indicated that an IgG-lambda antibody was being produced. At this stage, total RNA was prepared from approximately 10^6 cells. First strand cDNA synthesis was performed as described above using the primers HulgG1-4CG1FOR and HuCLFOR. Primary PCRs were performed for the VH-CH1 using a mixture of the 6 HuVHBACK2 primers and HulgG1-4CG1FOR and for the V lambda-C lambda using a mixture of the 7 HuV BACK primers and HuC FOR. Restriction, cloning and screening proceeded as described. To determine the diversity of the clones, the VH and V lambda genes of 15 clones were PCR amplified, restricted with the frequent cutting restriction enzyme BstNI and analyzed on a 4% agarose gel (see example 16).
- 25 30 35

- [0452]** Assay for anti-Rh-D activity and demonstration of specificity: A 5% (vol/vol) suspension of either Rh-D positive (OR2R2) or Rh-D negative (Orr) erythrocytes in phosphate buffered saline (PBS, pH 7.3) were incubated with a papain solution for 10 min at 37°C. The erythrocytes were washed three times in PBS and a 1% (vol/vol) suspension of erythrocytes was made up in PBS supplemented with 1% (vol/vol) of bovine serum albumin (BSA). Fifty ul of a papain treated erythrocyte suspension and 50ul of phage supernatant were placed in the wells of round bottom microtitre plates and the plates were placed on a Titertek plate shaker for 2 min. After 15 min incubation at 37°C 100 ul of PBS/BSA was added to each well. The plates were centrifuged at 200 g for 1 min and the supernatant was discarded. The erythrocytes were resuspended in the remaining PBS/BSA and the Fab fragments were crosslinked by addition of the 9E10 monoclonal antibody (50ul a 1ug/ml solution in PBS/BSA) directed against the myc peptide tag (Ward, E.S., et al., Nature 1989. supra). The plates were placed at room temperature (RT) until sedimentation had occurred. Agglutination of erythrocytes caused a diffuse button of erythrocytes and the results were evaluated macroscopically. Specificity was confirmed with a standard prepapainized (as above) panel of 9 erythrocyte suspensions in PBS (all suspensions blood group O, 4 D positive and 5 D negative) known to have homozygous expression of all the clinically relevant erythrocyte blood group alloantigens. The number of copies of the D antigen on the D positive cells varied between 10,000 and 20,000 per erythrocyte depending on the Rh genotype. Briefly, 50 ul phage supernatant in PBS supplemented with 2% (vol/vol) skimmed milk was mixed with 50 ul of a 2% erythrocyte suspension in PBS in glass tubes and incubated for 15 min at 37°C. After one wash with PBS/BSA the erythrocytes were pelleted and resuspended in 50 ul donkey anti-human lambda light chain (Sigma L9527, diluted 1:40 in PBS/BSA). The tubes were centrifuged for 1 min at 200g and agglutination was read macroscopically using "tip and roll" method.
- 40 45 50 55

R sults

[0453] a PCR assembly of a Fab from a human hybridoma: A single band of the correct size was obtained after amplification. Thirty-eight of 96 clones (40%) screened specifically agglutinated Rh-D positive but not Rh-D negative red blood cells. The results demonstrate a high frequency of successful splicing in the assembly process and the potential of this technique for one step cloning of human hybridomas.

[0454] b Assembly of human Fabs from a polyclonal lymphoblastic cell line (LCL): Analysis of the diversity of the clones indicated that 3 different heavy chain families and 2 different light chains families were present. Five anti-Rh-D specific clones were identified out of 96 screened. The VH and Vλ chains had identical nucleotide sequences in each clone and were typical of anti-Rh-D V-genes (unpublished results). The results demonstrate the potential of this technique to assemble, clone and isolate human antibody fragments from polyclonal cell populations (see also section on isolation of specific binding activities from an 'unimmunized' human library (examples 32 and 33).

Example 31

Selection of Phage Displaying a Human Fab Fragment directed against the Rhesus-D Antigen by binding to Cells displaying the Rhesus D Antigen on their Surface

[0455] A large number of important antigens are integral components of cell surface membranes, i.e. they are cell surface antigens. These include tumor specific antigens and red and white blood cell surface antigens. In many instances, it would be important to isolate antibodies against these antigens. For example, antibodies directed against the rhesus-D (Rh-D) antigen on red blood cells are used both diagnostically and therapeutically. Many of these antigens are difficult to purify and some, like Rh-D, are not biologically active when isolated from the membrane. Thus, it would be useful to be able to affinity purify antibody fragments displayed on the surface of bacteriophage directly on cell surface antigens. To test the feasibility of affinity purification on cell surface antigens, the anti-Rh-D human monoclonal antibody Fog-B was displayed as a Fab fragment on the surface of bacteriophage fd. The displayed Fog-B Fab fragment bound antigen as determined by agglutination assay and could be affinity purified on the basis of its binding on the surface of Rh-D positive red blood cells but not Rh-D negative red blood cells.

Materials and Methods

[0456] Construction of a clone encoding an anti-Rh-D Fab fragment in phagemid pHEN1 and display of the Fab fragment on the surface of bacteriophage fd.

[0457] The human hybridoma Fog-B has been previously described (N.C. Hughes-Jones et al Biochem, J. 268 135 (1990). It produces an IgG-1/lambda antibody which binds the Rh-D antigen. RNA was prepared from 10⁷ hybridoma cells using a modified method of Cathala (as described in example 12) and 1st strand cDNA synthesized using specific immunoglobulin heavy and light chain primers (HuVH1FOR [example 40] and HuCλ FOR (5' -GGA ATT CTT ATG AAG ATT CTG TAG GGG CCA C-3')) as described in example 14. The VH gene was subsequently amplified from an aliquot of the 1st strand cDNA using HuVH4aBACK and HuVH1FOR. The Vλ gene was amplified using a V primer specific for Fog-B (VλFog-B, 5'-AAC CAG CCA TGG CC AGT CTG TGT TGA CGC AGC C-3'). The PCR conditions were as described in example 40. The PCR products were analyzed by running 5μl on a 2% agarose gel. The remainder was extracted twice with ether, twice with phenol/chloroform, ethanol precipitated and resuspended in 50μl of H₂O. The amplified VH DNA was digested with Pst1 and BstEII, and the amplified Vλ-Cλ DNA with NcoI and EcoR1. The fragments were purified on a 2% agarose gel, extracted using GeneClean, and sequentially ligated into the soluble expression vector pJM-1 Fab D1.3 (Fig 38). Clones containing the correct insert were initially identified by restriction analysis and verified by assay of expressed soluble Fab (see example 19 for induction conditions). The Fog-B Fab cassette was amplified from pJM-1 by PCR using HUVH4BACK-Sfi and Hu Cλ-Not, digested with the appropriate restriction enzymes and ligated into pHEN1. Clones containing the correct insert were identified initially by restriction analysis and subsequently by assay (see example 21 for induction conditions).

[0458] Assay for soluble Fog-B Fab fragment and phage displayed Fog-B Fab fragment for anti-Rh-D activity and documentation of specificity.

[0459] Assay of the soluble expressed Fab was performed on unconcentrated E.coli supernatant. Assay of Fog-B displayed on the phage surface was performed on phage that had been concentrated 10 fold by PEG precipitation and then resuspended in PBS. the assays for activity and specificity are as described in example.

[0460] Cell surface affinity purification of phage displaying Fog-B anti-Rh-D Fab fragment

[0461] Purified Fog-B phage was mixed with purified phage Fd-Tet CAT-1 displaying the anti-lysozyme scFv D1.3 (pAbD1.3) in a ratio of approximately 1 Fog-B:50 scFvD1.3. Prepapainized erythrocytes (OR2R2 [Rhesus positive] or Orr [Rhesus negative]) were suspended in PBS supplemented with 2% skimmed milk powder in a concentration of

4x10⁷/ml. One ml of this suspension was mixed with 10¹¹ phage suspended in 2 ml of PBS supplemented with 2% skimmed milk and incubated for 30 min at room temperature under continuous rotation. The erythrocytes were washed three times with an excess of ice-cold PBS (10 ml per wash) and subsequently pelleted. The phage were eluted from the cells by resuspending in 200 µl of 76 mM citric acid pH 2.8 in PBS for 1 min. The cells were then pelleted by centrifugation for 1 min at 3000 rpm and the supernatant containing the eluted phage was neutralized by adding 200 µl of 240 mM Tris-base, 22mM Disodium hydrogen phosphate in 1% w/vol albumin. Serial dilutions of the eluate was used to infect TG1 cells. Fog-B Fab phage were selected on ampicillin plates and scFvD1.3 phage on tetracycline plates and the titre of each determined prior to selection, after selection on rhesus-D negative cells and after selection on rhesus-D positive cells.

Results

[0462] Fog-B Fab fragment displayed on the surface of the phage derived from the phagemid pHEN clone specifically agglutinated rhesus-D positive but not rhesus D-negative red blood cells. Affinity purification of the Fog-1 Fab phagemid on Rh-D positive red blood cells resulted in an enrichment from 1:50 to 1500:1 (Fog-B Fab:scFvD1.3), whereas purification on Rh-D negative red blood cells demonstrated essentially no enrichment (10 fold).

	TITRE	RATIO
	Fog-B Fab scFvD1.3	Fog-B Fab/scFvD1.3
Prior to selection	1.0 x 10 ⁹ 5.0 x 10 ⁹	1:50
Selection on Rh-D negative cells	2.0 x 10 ⁴ 1.0 x 10 ⁵	1:5
Selection on Rh-D positive cells	6.0 x 10 ⁶ 4.0 x 10 ³	1500:1

Example 32 A PCR Based Technique for One Step Cloning of Human scFv Constructs

[0463] Assembly of human scFv is similar to the assembly of mouse scFvs described in example 12. To develop the PCR cloning of human V genes it was necessary to design a new range of human specific oligonucleotide primers (table 60). The use of these primers for the generation of human Fabs is described in example 30. The assembly of human scFvs is essentially the same but requires a set of FORWARD primers complementary to the J segments of the VH, VK and V lambda genes. (For Fabs FORWARD primers complementary to the constant region are used.) The J segment specific primers were designed based on the published JH, JK and J lambda sequences (Kabat, E.A. et al; Sequences of Proteins of Immunological Interest. 4th Edition. US Department of Health and Human Services. 1987).

[0464] In addition, a different linker is needed for scFvs than for Fabs so for human scFvs a new set of primers was needed to prepare the linker. Primers complementary to the JH forward primers and the VK and V lambda back primers were synthesized to permit generation of linker DNA by PCR amplification of a plasmid template containing the scFv linker (Table 6, Fig. 39). To ensure adequate amplification, the primers were extended into the actual linker sequence. Using these primers to make the scFv linker DNA, 52 separate PCR reactions were performed using each of the 4 reverse JH primers in combination with each of the 13 reverse VK and V lambda oligonucleotides. The template was approximately 1ng of pSW2scD1.3 (Ward, E.S. 1989 supra) containing the short peptide (Gly4Ser)3 "(Huston, J.S. et al., Gene 1989. 77:61)

A specific example of PCR assembly of a human scFv library

[0465] This example describes the generation of a human library of scFvs made from an unimmunized human:

[0466] 500ml of blood, containing approximately 10⁸ B-cells, was obtained from a healthy volunteer blood donor. The white cells were separated on Ficoll and RNA was prepared as described in example 12.

[0467] Twenty percent of the RNA, containing the genetic material from approximately 2 x 10⁷ B-cells, was used for cDNA preparation as described in example 30. Heavy chains originating from IgG and IgM antibodies were kept separate by priming cDNA synthesis with either an IgG specific primer (HulgG1-4CH1FOR) or an IgM specific primer (HulgMFOR). Aliquots of the cDNA was used to generate four separate scFv libraries (IgG-K, IgG-lambda, IgM-K and IgM-lambda) as described in example 30. The resulting libraries were purified on 1.5% agarose, electroeluted and ethanol precipitated. For subsequent cloning, the K and lambda libraries were combined giving separate IgG and IgM libraries.

[0468] Cloning of the library: The purified scFv fragments (1-4ug) were digested with the restriction enzymes NotI and either SfiI or NcoI. After digestion, the fragments were extracted with phenol/chloroform, ethanol precipitated. The digested fragments were ligated into either SfiI-NotI or NcoI-NotI digested, agarose gel electrophoresis purified pHEN1

DNA (6 μ g) (see example 20), in a 100 μ l ligation mix with 2,000 U T4 DNA ligase (New England Biolabs) overnight at room temperature. The ligation mix was purified by phenol extraction and ethanol precipitated. The ligated DNA was resuspended in 10 μ l of water, and 2.5 μ l samples were electroporated into E.coli TG1 (50 μ l). Cells were grown in 1 ml SOC for 1 hr and then plated on 2 x TY medium with 100 μ g/ml ampicillin and 1% glucose (AMP-GLU), in 243 x 243 mm dishes (Nunc). After overnight growth colonies were scraped off the plates into 10 ml 2 x TY containing AMP-GLU and 15% glycerol for storage at -70°C as a library stock.

[0469] Cloning into Sfil-NotI and NcoI-NotI digested pHEN1 yielded libraries of 10^7 and 2×10^7 clones respectively for the IgM libraries and approximately 5×10^7 clones for each of the two IgG libraries.

Example 33 Isolation of binding activities from a library of scFvs from an unimmunized human

[0470] The ability to select binding activities from human antibody libraries displayed on the surface of phage should prove even more important than isolation of binding activities from murine libraries. This is because the standard way of generating antibodies via hybridoma technology has not had the success with human antibodies that has been achieved with mouse. While in some instances it will be possible to make libraries from immunized humans, in many cases, it will not prove possible to immunize due to toxicity or lack of availability of an appropriate immunogen or ethical considerations. Alternatively, binding activities could be isolated from libraries made from individuals with diseases in which therapeutic antibodies are generated by the immune response. However, in many cases, the antibody producing cells will be located in the spleen and not available in the circulating pool of peripheral blood lymphocytes (the most easily accessible material for generating the library). In addition, in diseases associated with immunosuppression, therapeutic antibodies may not be produced.

[0471] An alternative approach would be to isolate binding activities from a library made from an unimmunized individual. This approach is based on estimates that a primary repertoire of 10^7 different antibodies is likely to recognize over 99% of epitopes with an affinity constant of 10^5 M⁻¹ or better. (Pewrelson, A.S. Immunol. Rev. (1989) 110:5). While this may not produce high affinity antibodies, affinity could be boosted by mutation of the V-genes and/or by using the isolated VH domain in a hierarchical approach with a library of light chains (or vice versa). In this section, we demonstrate the feasibility of this approach by isolating specific antigen binding activities against three different antigens from a library of scFvs from an unimmunized human.

Materials and Methods

[0472] The generation of the human scFv library used for the isolation of binding activities described in this example is detailed in example 32.

[0473] Estimation of diversity of original and selected libraries: Recombinant clones were screened before and after selection by PCR (example 16) with primers LMB3 (which sits 5' of the pelB leader sequence and is identical to the reverse sequencing primer (-40 n) of pUC19) and fd-SEQ1 followed by digestion with the frequent-cutting enzyme BstNI. Analysis of 48 clones from each unselected library indicated that 90% of the clones had inset, and the libraries appeared to be extremely diverse as judged by the BstNI restriction pattern.

Rescue of Phagemid libraries for enrichment experiments:

[0474] To rescue phagemid particles from the library, 100 ml 2 x TY containing AMP-GLU (see example 32) was inoculated with 10^9 bacteria taken from the library (prepared in example 42) (approx. 10 μ l) and grown for 1.5 hr, shaking at 37°C. Cells were spun down (IEC- centrifuge, 4 K, 15 min) and resuspended in 100 ml prewarmed (37°C) 2 x TY-AMP (see example 31) medium, 2×10^{10} pfu of VCS-M13 (Stratagene) particles added and incubated 30 min at 37° without shaking. Cells were then transferred to 900 ml 2 x TY containing ampicillin (100 μ g/ml) and kanamycin (25 μ g/ml) (AMP-KAN), and grown overnight, while shaking at 37°C. Phage particles were purified and concentrated by three PEG-precipitations (see materials and methods) and resuspended in PBS to 10^{13} TU/ml (ampicillin resistant clones).

Enrichment for phOx:BSA binders by selection on tubes:

[0475] For enrichment, a 75 x 12 mm Nunc-immunotube (Maxisorp; Cat. No. 4-44202) was coated with 4 ml phOx:BSA (1 mg/ml; 14 phox per BSA in 50 mM NaHCO3 pH 9.6 buffer) overnight at room temperature. After washing three times with PBS, the tube was incubated for 2 hr at 37°C with PBS containing 2% Marvel (2% MPBS) for blocking. Following three PBS washes, phagemid particles (10^{13} TU) in 4 ml of 2% MPBS were added, incubated 30 min at room temperature on a rotating turntable and left for a further 1.5 hours. Tubes were then washed with 20 washes of PBS, 0.1% Tween 20 and 20 washes PBS (each washing step was performed by pouring buffer in and out immediately).

Bound phage particles were eluted from the tube by adding 1 ml 100 mM triethylamine pH 11.5 and rotating for 15 min. The eluted material was immediately neutralised by adding 0.5 ml 1.0 M Tris-HCl, pH 7.4 and vortexed. Phage was stored at 4°C.

[0476] Eluted phage (in 1.5 ml) was used to infect 8 ml logarithmic growing E.coli TG1 cells in 15-ml 2 x TY medium, and plated on AMP-GLU plates as above yielding on average 10^7 phage infected colonies.

[0477] For selection of phOx:BSA binders, the rescue-tube enrichment -plating cycle was repeated 4 times, after which phagemid clones were analysed for binding by ELISA. Enrichment for lysozyme binders by panning and on columns: A petri dish (35 x 10 mm Falcon 3001 Tissue culture dish) was used for enrichment by panning. During all steps, the plates were rocked on an A600 rocking plate (Raven Scientific). Plates were coated overnight with 1 ml turkey egg white lysozyme (3 mg/ml) in 50 mM sodium hydrogen carbonate (pH 9.6), washed three times with 2 ml PBS, and blocked with 2 ml 2% MPBS at room temperature for 2. hours. After three PBS washes approximately 10^{12} TU phage particles in 1 ml 2% MPBS were added per plate, and left rocking for 2 hr at room temperature. Plates were washed for 5 min with 2 ml of the following solutions: 5 times PBS, PBS-Tween (0.02% Tween-20), 50 mM Tris-HCl (pH 7.5) + 500 mM NaCl, 50 mM Tris-HCl (pH 8.5) + 500 mM NaCl, 50 mM Tris-HCl (pH 9.5) + 500 mM NaCl and finally 50 mM sodium hydrogen carbonate pH 9.6 Bound phage particles were then eluted by adding 1 ml 100 mM triethylamine pH 11.5 and rocking for 5 min before neutralising with 1 M Tris-HCl (pH 7.4) (as above). Alternatively, 1 ml turkey egg white lysozyme-Sepharose columns were used for affinity purification (McCafferty, J., et al., Nature 1990. 348: 552) Columns were washed extensively with PBS, blocked with 15 ml 2% MPBS, and phage (10^{12} TU) in 1 ml 2% MPBS loaded. After washing with 50 ml PBS, 10 ml PBS- Tween (PBS - 0.02% Tween-20), 5 ml of 50 mM Tris-HCl (pH 7.5) - 500 mM NaCl, 5 mM Tris-HCl 9pH 8.5) + 500 mM NaCl, 5ml of 50 mM Tris-HCl (pH 9.5) + 500 mM NaCl and finally 5 ml of 50 mM sodium hydrogen carbonate pH 9.6. Bound phage was eluted using 1.5 ml 100 mM triethylamine and neutralised with 1 M Tris-HCl (pH 7.4).

[0478] For selection of turkey egg white lysozyme binders, the rescue-tube enrichment-plating cycle or rescuolumn-plating cycle was repeated 4 times, after which phagemid clones were analysed for binding by ELISA. Rescue of individual phagemid clones for ELISA: Clones resulting from reinfecting and plated phage particles eluted after 4 rounds of enrichment, were inoculated into 150 µl of 2 x TY-AMP-GLU in 96-well plates (cell wells, Nunc), grown with shaking (250rpm) overnight at 37°C. A 96-well plate replicator ('plunger') was used to inoculate approximately 4 µl of the overnight cultures on the master plate into 200 µl fresh 2 x TY-AMP-GLU. After 1 hr, 50 µl 2 x TY-AMP-GLU containing 10^8 pfu of VCS-M13 was added to each well, and the plate incubated at 37°C for 45 min, followed by shaking the plate at 37°C for 1 hr. Glucose was then removed by spinning down the cells (4K, 15 min), and aspirating the supernatant with a drawn out glass pasteur pipet. Cells were resuspended in 200 µl 2 x TY-AMP-KAN (Kanamycin 50 µg/ml) and grown 20 hr, shaking 37°C. Unconcentrated supernatant containing phage was taken for analysis by ELISA.

ELISA

[0479] Analysis for binding to phOx:BSA, BSA or lysozyme was performed by ELISA (see example 9), with 100 µg/ml phOx:BSA or BSA, or 3 mg/ml turkey egg white lysozyme used for coating. Determination of cross reactivity to unrelated antigens with the isolated clones was also determined by ELISA on plates coated with 100 µg/ml of an irrelevant antigen (keyhole limpet haemocyanin (KLH), ovalbumin, chymotrypsinogen, cytochrome C, thyroglobulin, GAP-DH (glyceraldehyde-3-phosphate dehydrogenase), or trypsin inhibitor). Characterization of ELISA positive clones: All antigen specific clones isolated were checked for cross reactivity against a panel of irrelevant antigens as described above. The diversity of the clones was determined by PCR screening as described above and at least two clones from each restriction pattern were sequenced by the dideoxy chain termination method.

Results

[0480] Isolation and characterization of phOx:BSA binders: After 4 rounds of selection, ELISA-positive clones were isolated for phOx:BSA. All clones originated from the IgM library. Of 96 clones analysed, 43 clones were binding to both phOx:BSA and BSA, with ODs ranging from 0.4 to 1.3 (background 0.125). These clones are designated as BSA binders. The binding to BSA seemed to be specific, since none of the 11 clones analysed gave a signal above background when used in an ELISA with KLH, ovalbumin, chymotrypsinogen, cytochrome C, lysozyme, thyroglobulin, GAP-DH, or trypsin inhibitor, all BSA binding clones had the same BstNI restriction pattern, and 14 clones were completely sequenced. Thirteen of the fourteen clones had the same sequence, the VH was derived from a human VH3 family gene and the VL from a human V lambda 3 family gene (Table 1). The other BSA binder was derived from a human VH4 family gene and a human Vk1 family gene (data not shown).

[0481] One clone was isolated which bound to phox:BSA only (OD 0.3), and bound phage could be completed off completely by adding 0.02 mM 4-E-amino-caproic acid methylene 2-phenyl-oxazol-5-one (phOx-CAP) as a competitor. Also no binding above background could be detected to the panel of irrelevant proteins described above. The sequence

revealed a VH derived from a human VH1 family gene and a VL derived from a human V lambda 1 family gene (Table 7).

Isolation and characterisation of lysozyme binders:

5 [0482] After 4 rounds of selection, 50 ELISA-positive clones were isolated for turkey lysozyme. The majority of the clones, greater than 95%, were from the IgM library. The binding to lysozyme seemed to be specific, since none of the clones analysed gave a signal above background when used in an ELISA with KLH, ovalbumin, chymotrypsinogen, cytochrome C, thyroglobulin, GAP-DH, or trypsin inhibitor. The lysozyme binding clones gave 3 different BstNI restriction patterns, and at least 2 clones from each restriction pattern were completely sequenced. The sequences indicated
10 the presence of 4 unique human VH-VL combinations. (Table 7).

Conclusion

15 [0483] The results indicate that antigen binding activities can be isolated from repertoires of scFvs prepared from IgM cDNA from human volunteers that have not been specifically immunized.

Example 34

Rescue of human IgM library using helper phage lacking gene 3 ($\delta g3$)

20 [0484] This example describes the rescue of gene 3 fusions from a human library using a helper phage with a gene 3 deletion.

[0485] 100 μ l of bacterial stock of the IgM phagemid library prepared as described (example 32), containing 5×10^8 bacteria, was used to inoculate 100mls of 2xTY medium containing 100 μ g/ml ampicillin, 2% glucose (TY/Amp/Glu).
25 This was grown at 37°C for 2.5 hours. 10 mls of this culture was added to 90 mls of prewarmed TY/Amp/Glu and infection carried out by adding 10mls of a 200 fold concentrate of K07 helper phage lacking gene 3 (M13KO7gIII Δ No. 3) (example 27) and incubating for 1 hour at 37°C without shaking. Preparation of M13KO7gIII No.3 was as described in example 27. After centrifugation at 4,000 r.p.m. for 10 minutes the bacteria were resuspended in 100 ml of 2 x TY medium containing 100 μ g/ml ampicillin (with no glucose). Titration of the culture at this point revealed that there were
30 1.9×10^8 infected bacteria as judged by their ability to grow on plates containing both ampicillin (100 μ g/ml) and kanamycin (50 μ g/ml). Incubation was continued for 1 hour with shaking before transferring to 2.5 litres of 2xTY medium containing 100 μ g/ml ampicillin, 50 μ g/ml kanamycin, contained in five 2.5 litre flasks. This culture was incubated for 16 hours and the supernatant prepared by centrifugation. (10-15 minutes at 10,000 r.p.m. in a Sorvall RCSB centrifuge at 4°C). Phage particles were harvested by adding 1/5th volume of 20% polyethylene glycol, 2.5 M NaCl, standing at
35 4°C for 30 minutes and centrifuging as above. The resulting pellet was resuspended in 40mls of 10mM Tris, 0.1mM EDTA pH 7.4 and bacterial debris removed by centrifugation as above. The packaged phagemid preparation was then re-precipitated, collected as above and resuspended in 10mls of 10mM Tris, 0.1mM EDTA pH 7.4. The titre of this preparation was 4.1×10^{13} transducing units/ml (ampicillin resistance).

[0486] Tubes coated with OX-BSA were prepared as described in example 35 for panning the phagemid library from
40 example 32. The rescued library was also panned against tubes coated with bovine thyroglobulin (Sigma). These were coated at a concentration of 1mg/ml thyroglobulin in 50mM NaHCO₃ pH9.6 at 37°C, overnight. Tubes were blocked with PBS containing 2% milk powder (PBS/M) and incubated with 1ml of the rescued phagemid library (the equivalent of 250mls of culture supernatant) mixed with 3mls of PBS/M for 3 hours. Washing, elution, neutralisation and infection were as described in example 35.

45 Results: Panning against oxazalone - BSA

[0487] The first round of panning against OX-BSA yielded 2.8×10^6 phage. A large bacterial plate with 1.4×10^6 colonies derived from this eluate was scraped into 10mls of 2xTY, 20% glycerol, shaken for 10 minutes, aliquoted and stored.
50 This was also used to inoculate a fresh culture for rescue with M13KO7gIII Δ No.3. (Bacteria and rescued phage derived from first round panning against OX-BSA are named OXPAN1. Bacteria or rescued phage derived from second and third round panninas are named OXPAN2 and OXPAN3 respectively) Rescue of phagemid with M13KO7gIII Δ No.3 after each round of panning was essentially as described above but using 5ml volumes for the initial cultures in TY/Amp/Glu, using 1ml of helper phage and transferring to 100-500mls of 2xTY medium containing 100 μ g/ml ampicillin,
55 50 μ g/ml kanamycin. Second and third round panning steps were as described above for the first round, but using 0.8-1.0mls of 100 fold concentrated phage (the equivalent of 80-100 mls of culture supernatant). The eluate from the second round panning contained 8×10^8 infectious particles and the eluate from the third round panning contained 3.3×10^9 infectious particles.

Panning against thyroglobulin

[0488] The first round panning against thyroglobulin yielded 2.52×10^5 infectious particles. Half of the eluate was used to generate 1.26×10^5 bacterial colonies on a large plate. These colonies were scraped into 10mls of 2xTY, 20% glycerol, shaken for 10 minutes, aliquoted and stored. These bacteria and rescued phage derived from them are termed THYPAN1, and used to inoculate a fresh culture for rescue with M13KO7gIII No.3 to give a polyclonal rescued phage preparation. Material similarly derived from second and third round pannings are termed THYPAN2 and THYPAN3 respectively. Second and their round pannings with thyroglobulin were as described for second and third round OX-BSA panning. The eluate from the second round panning contained 8×10^7 transducing units and the eluate from the third round panning contained 6×10^7 infectious particles.

ELISA screening of clones derived by panning

[0489] 40 colonies derived from the third round of panning against thyroglobulin (THYPAN3) were picked into a 96 well plate and grown overnight at 37°C in 200µl of TY/Amp/Glu. Similarly 48 colonies from two rounds and 48 colonies from three rounds of panning against OX-BSA were grown (OX-PAN2 and OX-PAN3). Polyclonal phage were prepared at the same time. Next day 5µl from each culture was transferred to 100µl of fresh prewarmed TY/Amp/Glu grown for 1.5 hours and M13KO7gIII No.3 added (2×10^5 infectious phage per well in 100µl of TY/Amp/Glu). these were incubated for 1 hour at 37°C without shaking, centrifuged at 4,000 r.p.m. for 10 minutes, resuspended in 150µl of 2xTY medium containing 100µg/ml ampicillin and incubated for a further hour with shaking before adding to 2mls of medium containing 100µg/ml ampicillin, 50µg/ml kanamycin. After overnight growth the cultures were centrifuged at 4,000 r.p.m. for 10 minutes and the supernatants collected. ELISA plates used to screen THYPAN3 clones were coated at 37°C overnight with 200µg/ml thyroglobulin in 50mM NaHCO₃pH9.6. Plates used for OXPAN2 and OXPAN3 were coated at 100µg/ml OX-BSA in PBS at 37°C overnight.

[0490] 120µl of culture supernatant was mixed with 30ul of 5x PBS, 10% milk powder and incubated at room temperature for 2 hours at room temperature. ELISAs were carried out as described in example 14.

[0491] For thyroglobulin, 18 out of 40 clones were positive (0.3-2.0 O.D. after 30 minutes). (A phage control (vector pCAT3) gave a reading of 0.07 O.D.). In addition, positives were also seen on the polyclonal phage preparations THYPAN1 (0.314 O.D.) and THYPAN2 (0.189 O.D.) compared with phage derived from the original non-panned phagemid library (0.069 O.D.). All polyclonal phage were PEG precipitated and used at a 10 fold concentration.

[0492] PCR reactions and BstNI digests were carried out on the positive clones as described above and six different patterns of DNA fragments were obtained showing that at least six different clones had been isolated.

[0493] For OX-BSA after two rounds of panning, 30 of 48 clones were positive by ELISA and after three rounds, 42 of 48 were positive. In a separate experiment, positive signal was obtained from the polyclonal phage preparations OXPAN1 (0.988 OD) and OXPAN2 (1.717 OD) compared with phage derived from the original non-panned phagemid library (0.186 O.D.) after 30 minutes.

Specificity of clones for thyroglobulin or OX-BSA

[0494] Selected clones (11 anti-thyroglobulin, 5 anti-OX-BSA) representing each of the different BstNI restriction digest patterns were assayed for binding to a panel of irrelevant antigens. ELISA plates were coated with antigen (100 µl/ml in 50 mM NaHCO₃, pH 9.6) by overnight incubation at 37°C. The panel of antigens consisted of keyhole limpet haemocyanin, hen egg lysozyme, bovine serum albumin, ovalbumin, cytochrome c, chymotrypsinogen, trypsin inhibitor, GAP-D11 (glyceraldehyde-3-phosphate dehydrogenase), bovine thyroglobulin and oxazolone-BSA. Duplicate samples of phage supernatant (80 µl + 20 µl 5 x PBS, 10% milk powder) were added to each antigen and incubated for 1 hour at room temperature. the ELISA was carried out as described in example 14.

[0495] Each of the thyroglobulin specific clones (11 from 11) were positive for thyroglobulin (OD 0.12 - 0.76) but after 60 minutes showed no binding (OD<0.03) to any of the 9 irrelevant antigens. Similarly of the 5 OX-BSA specific clones 3 had an OD 0.07 - 0.52 compared to ODs < 0.02 for the irrelevant antigens. None of the 5 clones had any binding to BSA alone.

[0496] Thus positive clones can be isolated after only two rounds of panning by rescuing with M13KO7gIII No.3. In addition there is a greater likelihood with this helper of generating phage particles with more than one intact antibody molecule. This will potentially increase the avidity of phage-antibodies and may enable isolation of clones of weaker affinity.

Example 35: Alteration of fine specificity of scFv D1.3 displayed on phage by mutagenesis and selection on immobilised turkey lysozyme

[0497] The D1.3 antibody binds hen egg lysozyme (HEL) with an affinity constant of $4.5 \times 10^6 \text{M}^{-1}$ whereas it binds turkey egg lysozyme (TEL) with an affinity of $<1 \times 10^5 \text{M}^{-1}$. (Harper et al (1987) *Molecular Immunology* 24 p97-108, Amit et al (1986) *Science* 233 p747-753).

[0498] It has been suggested that this is because the glutamine residue present at position 121 of HEL (gln121) is represented by histidine residue at the same position in TEL. Thus mutagenising the D1.3 antibody residues which interact with gln121 of HEL may facilitate binding to TEL.

[0499] According to Amit et al, supra, tyrosine at amino acid position 32, phenylalanine at position 91 and tryptophan at position 92 of the light chain interact with gln121 of HEL. In addition tyrosine at position 101 of the heavy chain also interacts. None of these residues are predicted to be involved in determining the main chain conformation of the antibody variable regions (Chothia and Lesk (1987) *Journal of Molecular Biology* 196, p901-917).

Mutagenesis of pCAT3SCFvD1.3

[0500] The oligonucleotides mutL91,92, was prepared to randomise phenylalanine at position 91 (L91) and tryptophan at position 92 (L92) of the light chain. The oligonucleotides mutL32, was prepared to randomise tyrosine at light chain position 32 (L32) and the oligonucleotides mutH101 was prepared to randomise tyrosine at position 101 of the heavy chain (H101). mutL91,92:

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5' CGT CCG AGG AGT ACT NNN NNN ATG TTG ACA GTA ATA 3'
mutL32:
5' CTG ATA CCA TGC TAA NNN ATT GTG ATT ATT CCC 3'
mutH101:
5' CCA GTA GTC AAG CCT NNN ATC TCT CTC TCT GGC 3'

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N represents a random insertion of equal amounts of A,C,G or T) in vitro mutagenesis of the phagemid vector, pCAT3scFvD1.3 (example 13) with the oligonucleotide mutL91,92 was carried out using an in vitro mutagenesis kit (Amersham). The resultant DNA was transformed by electroporation into TG1 cells using a Bio-Rad electroporator. 78,000 clones were obtained and these were scraped into 15mls of 2xTY/20% glycerol. This pool was called D1.3L91L92. Single stranded DNA was prepared by rescue with M13K07 as described in Sambrook et al, 1989 supra, and sequenced with the primer FDTSEQ1, using a Sequenase sequencing kit (United States Biochemical Corporation).

[0501] This revealed that the DNA had been successfully mutagenised as judged by the presence of bands in all four DNA sequencing tracks at the nucleotide positions encoding L91 and L92. This mutagenised single stranded DNA was subjected to a further round of mutagenesis as above using either mutL32 or mutH101 oligonucleotides. Mutagenesis with mutL32 gave rise to 71,000 clones (pool called D1.3L32) while mutH101 gave 102,000 clones (pool called D1.3H101). These clones were scraped into 15mls of 2xTY/20% glycerol. Single stranded DNA derived from each pool was sequenced with the oligonucleotides D1.3L40 and LINKSEQ1 respectively, as described above, and shown to be correctly randomised.

D1.3L40:

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5' CAG GAG CTG AGG AGA TTT TCC 3'
LINKSEQ1:
5' TCC GCC TGA ACC GCC TCC ACC 3'

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Preparation of rescued phage for affinity purification

[0502] 10-20µl of bacteria derived from each mutagenised pool (plate scrapes) was used to inoculate 5mls of TY/Glu/Amp. All bacterial growth was at 37°C. After 2-3 hours growth, 1ml was diluted in 5mls of prewarmed TY/Glu/Amp and infected by addition of 0.5 mls of a 200 fold concentrate of the M13K07gIII Δ No.3 preparation described in example 27. After 1 hour of infection the cultures were centrifuged at 4,000 r.p.m. for 10 minutes, resuspended in 2xTY, 100µg/ml ampicillin, incubated for a further hour, transferred to 500 mls of 2xTY medium containing 100 µg/ml ampicillin, 50 µg/ml kanamycin and grown for 16 hours. The remaining steps of phage preparation were as described in example 34. Phage were finally dissolved in 10mM Tris, 1mM EDTA pH7.4 at 1/100th the original culture volume.

Affinity purification

5 [0503] 10mls of turkey egg lysozyme at a concentration of 10mg/ml in 0.1M NaHCO₃, 0.5MNaCl pH8.3 was mixed with an equal volume of swollen Cyanogen Bromide Activated Sepharose 4B (Pharmacia), covalently linked and washed according to manufacturers instructions. Before use this matrix (TEL-Sepharose) was washed with 100 volumes of PBS followed by 10 volumes of PBSM. The TEL-Sepharose was resuspended in an equal volume of PBSM and 1ml was added to 1ml of a 50 fold concentrate of phage in PBSM and incubated on a rotating platform for 30 minutes at room temperature. The actual phage used for this step was prepared by mixing, equal volumes of the independent preparations of the three randomised pools (D1.3L9192, D1.3H101 and D1.3L32). After this binding step, 10 the suspensions were loaded onto a disposable polypropylene column (Poly-Prep columns, Bio-Rad) and washed with 200 volumes of PBS containing 0.1% Tween 20. Bound phage were eluted with 1ml of 100mM triethylamine and neutralised with 0.5ml 1M Tris (pH7.4). A dilution series was prepared from the eluate and used to infect TG1 cells and plated out on TY plates containing 100µg/ml ampicillin, 2% glucose. Plates carrying approximately 10⁶ colonies were scraped into 3mls of 2xTY, 20% glycerol and stored at -70°C. 10µl of this was used to initiate a second round 15 culture which was rescued with M13KO7gIIIΔ No.3 as described above (using a final culture volume of 100mls). Second and third round affinity column purification steps were carried out as described above for the first round.

Analysis by ELISA

20 [0504] 40 colonies derived from the third round of column purification on TEL-Sepharose were picked into a 96 well plate and grown overnight at 37°C in 200µl of TY/Amp/Glu. Phagemid particles were rescued and prepared for ELISA as described in example 18. ELISA plates were coated overnight at 37°C with hen egg lysozyme (HEL) or turkey egg lysozyme (TEL) at a concentration of 200µg/ml in 50mM NaHCO₃ pH9.6 ELISAs were carried out as described in example 14.

25 [0505] After 15 minutes incubation in substrate, 13 clones were found to be negative (OD<0.05 on HEL and TEL). In all positives, a signal of 0.1-0.78 was scored on HEL with the exception of one where signal on HEL was 0.078 but signal on TEL (OD 0.169) brought it in to the positive group. The control phagemid preparation had a percentage ratio of signal TEL:HEL of 22%. Clones were deemed to have an unaltered binding if the ratio of TEL:HEL was less than 40%. 9 clones fell into this category. 18 samples were scored as having altered binding with a ratio of signal on TEL: 30 HEL of between 40-200%.

[0506] A dilution series was made on 10 clones which were analysed by ELISA in 6 of these clones the profile of binding to HEL was the same as the original clone (pCAT3SCFvD1.3) while the signal with TEL was increased (see figure 50 clone B1). In the remaining 4 clones, the increased signal with TEL was accompanied by a decrease in signal on HEL (see figure 40 clone A4). 35

Competition with soluble antigen

[0507] All of the isolated clones retained binding to HEL to varying extents. In order to determine whether a soluble antigen could compete with the immobilised antigen, a parallel experiment was carried out, as above, but with the addition of hen egg lysozyme (1mg/ml) to TEL-Sepharose before incubating with the phage preparation. This experiment was carried through 3 rounds of column purification and 40 colonies were picked. None of these clones bound HEL or GEL demonstrating that the soluble antigen had been successful in competing out binding to the immobilised antigen. 40

45 Example 36Modification of the Specificity of an Antibody by Replacement of the VLK Domain by a VLK Library derived from an Unimmunised Mouse

50 [0508] When an antibody specificity is isolated it will often be desirable to alter some of its properties particularly its affinity or specificity. This example demonstrates that the specificity of an antibody can be altered by use of a different VL domain derived from a repertoire of such domains. This method using display on phage would be applicable to improvement of existing monoclonal antibodies as well as antibody specificities derived using phage antibodies. This example shows that replacement of the VL domain of scFvD1.3 specific for Hen eggwhite lysozyme (HEL) with a library 55 of VL domains allows selection of scFv fragments with bind also to Turkey eggwhite lysozyme (TEL). More generally this experimental approach shows that specificities of antibodies can be modified by replacement of a variable domain and gives a further example of the hierarchical approach to isolating antibody specificities.

[0509] The D1.3 heavy chain was amplified from an existing construct (pSW1-VHD1.3, Ward et al., 1989 supra) by

PCR using the primers VH1BACK and VH1FOR, the light chain library was amplified from a cDNA library derived from the spleen of an unimmunised mouse, which was synthesized by using the MJKFONX primers 1,2,4,5 for the first strand as in example 12. The subsequent amplification was performed with the same forward primers and the VK2BACK primer. The PCR assembly of the D1.3 heavy chain with the light chain library was mediated by the signal chain Fv linker as described in example 12.

[0510] Cloning the assembled PCR products (scFv sequences) was done after an additional PCR step (pull-through) using a BACK primer providing an ApaI site and forward primers which contained a Not I site as described in example 12. ApaI/Not I digested PCR fragments were cloned into the similarly digested vector fdCAT2. 5×10^5 transformations were obtained after electroporation of the ligation reaction into MC1061 cells.

[0511] Screening of the phage library for TEL binders was performed by panning. Polystyrene Falcon 2058 tubes were coated (16 hrs) with 2 ml of TEL-PBS (3 mg/ml) and blocked for 2 hrs with 4 ml MPBS (PBS containing 2% skimmed milk powder). Phage derived from the library (5×10^{10} transducing units) in 2 ml of MPBS (2%) were incubated in these tubes for 2 hrs at room temperature. The tubes were washed 3x with PBS, 1x with 50 mM Tris-HCl, pH 7.5, 0.5 M NaCl; 1x with 50mM Tris-HCl, pH8.5, 0.5 M NaCl, 50 mM Tris-HCl, pH 9.5 M NaCl. Finally phage were eluted with 100 mM triethylamine. Eluted phages were taken to infect TG1 cells, the cells were plated on 2xTY plates containing 15 µg/ml tetracycline and grown for 16h. The colonies were scraped into 25ml of 2xTy medium and the phages were recovered by PEG precipitation. After a second round of selection for TEL binders ELISAs were performed as described (example 2).

[0512] Analysis of 100 clones from the library before affinity selection by ELISA on plates coated with TEL showed no binders. In contrast, after two rounds of selection for TEL binding phages about 10% of the phage clones showed positive ELISA signals. ELISA signals were scored positive with values at least two fold higher than the fdCAT2 vector without insert. A more detailed analysis of binding properties of TEL binding phages is shown in figure 41.

[0513] As shown in figure 41, several clones were found which bind equally to TEL and HEL in contrast to the original D1.3 scFv, which binds almost exclusively to HEL. None of the clones bound to BSA. These findings indicate that the specificity of these scFvs was broader in comparison to D1.3, since both lysozymes (HEL and TEL) are recognized, but specificity for lysozyme was retained since other BSA was not recognized. The deduced amino acid sequences (derived by DNA sequencing) of two light chains from clones MF1 and M21, which correspond to clones 3 and 9 in figure 41 are shown in figure 42.

[0514] In the case of isolated antibodies the experimental approach as described in this study may be particularly useful if recognition of a wider range of different but closely related antigens is desired. For example, monoclonal antibodies against viral antigens like V3 loop of HIV-1 gp120 are in most cases quite specific for one particular virus isolate because of the variability in this part of the HIV-1 env gene. The modification of such antibodies in the way described in this example may lead to antibodies which cross react with a wider range of HIV-1 isolates, and would therefore be of potentially higher therapeutic or diagnostic value.

[0515] A similar approach could be taken in which a light chain variable domain of desired properties is kept fixed and combined with a library of heavy chain variable domains. Some heavy chains, for example VHD1.3 retain binding activity as single domains. This may allow a strategy where VH domains are screened for binding activity when expressed on phage and then binding domains combined with a library of VL domains for selection of suitable light chain partners.

Example 37

Selection of a Phage Antibody Specificity by Binding to an Antigen attached to Magnetic Beads. Use of a Cleavable Reagent to allow elution of Bound Phage under Mild Conditions

[0516] When a phage antibody binds to its antigen with high affinity or avidity it may not be possible to elute the phage antibody from an affinity matrix with a molecule related to the antigen. Alternatively, there may be no suitable specific eluting molecule that can be prepared in sufficiently high concentration. In these cases it is necessary to use an elution method which is not specific to the antigen-antibody complex. Unfortunately, some of the non-specific elution methods disrupt phage structure, for instance phage viability is reduced with time at pH12 (Rossomando, E.F. and Zinder, N.D. J. Mol. Biol. 36 387-399 1968). A method was therefore devised which allows elution of bound phage antibodies under -mild conditions (reduction of a dithiol group with dithiothreitol) which do not disrupt phage structure.

[0517] Target antigen was biotinylated using a cleavable biotinylation reagent. BSA conjugated with 2-phenyl-5-oxazolone (O. Makela et al. supra) was modified using a biotinylation reagent with a cleavable dithiol group (sulphosuccinimidyl 2-(biotinamido) ethyl-1,3-dithiopropionate from Pierce) according to the manufacturers instructions. This biotinylated antigen was bound to streptavidin coated magnetic beads and the complex used to bind phage. Streptavidin coated magnetic beads (Dynal) were precoated with antigen by mixing 650µg of biotinylated OX-BSA in 1 ml PBS, with 200µl of beads for at least 1 hour at room temperature. Free antigen was removed by washing in PBS. One fortieth

of the complex (equivalent to 5µl of beads and an input of 17.5 µg of OX-BSA) was added to 0.5ml of phage in PBSM (PBS containing 2% skimmed milk powder) containing 1.9×10^{10} phage particles mixed at the ratios of pAbD1.3 directed against lysozyme (example 2) to pAbNQ11 directed against 2-phenyl-5-oxazolone shown in Table 8.

[0518] After 1 hour of incubation with mixing at room temperature, magnetic beads were recovered using a Dynal MPC-E magnetic separation device. They were then washed in PBS containing 0.5% Tween 20, (3x10 minutes, 2x1 hour, 2x 10 minutes) and phage eluted by 5 minutes incubation in 50ul PBS containing 10mM dithiothreitol. The eluate was used to infect TG1 cells and the resulting colonies probed with the oligo NQ11CDR3

(5' AAACCAGGCCCGTAATCATAGCC 3') derived from CDR3 of the NQ11 antibody (This hybridises to pAbNO11 but not pAb D1.3).

[0519] A 670 fold enrichment of pAbNQ11 (table 8) was achieved from a background of pAbD1.3 in a single round of purification using the equivalent of 17.5µg of biotinylated OX-BSA.

[0520] This elution procedure is just one example of an elution procedure under mild conditions. A particularly advantageous method would be to introduce a nucleotide sequence encoding amino acids constituting a recognition site for cleavage by a highly specific protease between the foreign gene inserted, in this instance a gene for an antibody fragment, and the sequence of the remainder of gene III. Examples of such highly specific proteases are Factor X and thrombin. After binding of the phage to an affinity matrix and elution to remove non-specific binding phage and weak binding phage, the strongly bound phage would be removed by washing the column with protease under conditions suitable for digestion at the cleavage site. This would cleave the antibody fragment from the phage particle eluting the phage. These phage would be expected to be infective since the only protease site should be the one specifically introduced. Strongly binding phage could then be recovered by infecting e.g. E.coli TG1 cells.

Example 38

Use of Cell Selection to provide an Enriched Pool of Antigen Specific Antibody Genes. Application to reducing the Complexity of Repertoires of Antibody Fragment displayed on the Surface of Bacteriophage

[0521] There are approximately 10^{14} different combinations of heavy and light chains derived from the spleen of an immunised mouse. If the random combinatorial approach is used to clone heavy and light chain fragments into a single vector to display scFv, Fv or Fab fragments on phage, it is not a practical proposition to display all 10^{14} combinations. One approach, described in this example, to reducing the complexity is to clone genes only from antigen selected cells. (An alternative approach, which copes with the complexity is the dual combinatorial library described in example 22).

[0522] The immune system uses the binding of antigen by surface immunoglobulin to select the population of cells that respond to produce specific antibody. This approach of selecting antigen binding cells has been investigated to reduce the number of combinatorial possibilities and so increase the chance of recovering the original combination of heavy and light chains.

[0523] The immunological response to the hapten 4-hydroxy-3-nitrophenylacetic acid (NP) has been extensively studied. Since the primary immune response to NP uses only a single light chain the applicants were able to examine the use of the combinatorial method using a fixed light chain and a library of heavy chains to examine the frequencies genes that code for antibodies binding to NIP (4-hydroxy-3-iodo-5-nitrophenylacetic acid). The applicants have thus used this system to investigate the merits of selecting cell populations prior to making combinatorial libraries for display on phage.

Methods

2.1 Hapten conjugates

[0524] Chick gamma globulin (CGG, Sigma. Poole, UK) and Bovine serum albumen (BSA, Boehringer. Mannheim. Germany) were conjugated with NP-O-succinimide or NIP-caproate-O-succinimide (Cambridge Research Biochemicals, Northwich, UK) based on the method described by Brownstone (Brownstone, A., Mitchison, N.A. and Pitt-Rivers, R., Immunology 1966. 10: 465-492). The activated compounds were dissolved in dimethylformamide and added to proteins in 0.2 M sodium hydrogen carbonate. They were mixed with constant agitation for 16 hours at 4°C and then dialysed against several changes of 0.2 M sodium hydrogen carbonate. They were finally dialysed into phosphate buffered saline (PBS). The conjugates made were NP₁₂CGG, NIP₁₀BSA. The NIP₁₀BSA derivative was subsequently biotinylated using a biotinylation kit purchased from Amersham (Amersham International, Amersham, UK).

2.2 Animals and immunisation

[0525] Mice of the strain C57BL/6 were immunised by intraperitoneal injection of 100µg NP-CGG in Complete Fre-

unds Adjuvant at 10 weeks of age.

2.3 Spleen preparation

[0526] Seven days after immunization cells from the spleen were prepared as described by Galfre and Milstein (Galfre, G. and Milstein, C. *Methods Enzymol.* 1981. 73:3-46). Red cells were lysed with ammonium chloride (Boyle, W. *Transplantation* 1968.6:71) and when cell selection was performed dead cells were removed by the method described by von Boehmer and Shortman (von Boehmer, H. and Shortman, K, J. *Immunol, Methods* 1973.1:273). The cells were suspended in phosphage buffered saline (PBS), 1% Bovine serum albumen, 0.01% sodium azide; throughout all cell selection procedures the cells were kept at 4°C in this medium.

2.4 Cell Solution

[0527] Biotinylated NIP-BSA was coupled to streptavidin coupled magnetic beads (Dynabeads M280 Streptavidin, Dynal, Oslo, Norway) by incubating 10^8 beads with 100µg of biotinylated protein for 1 hour, with occasional agitation, and then washing five times to remove unbound antigen. The coupled beads were stored at 4°C in medium until required. For selection of antigen binding cells the cells (2×10^7 /ml) were first incubated for 30 minutes with uncoupled beads, at a bead: cell ratio of 1:1, to examine the degree of non-specific binding. The beads were then separated by placing the tube in a magnetic device (MPC-E Dynal) for 3-5 minutes. The unbound cells were removed and then incubated with NIP-BSA coupled magnetic beads, at a bead:cell ratio of 0.1:1, for 60 minutes, with occasional agitation. The beads and rosetted cells were separated as described above. The beads were then resuspended in 1 ml of medium and the separation repeated: this process was repeated 5-7 times until no unbound cells could be detected when counted on a haemocytometer.

[0528] For the depletion of surface immunoglobulin positive cells the cells were incubated with 20µg biotinylated goat anti-mouse polyvalent immunoglobulin (Sigma, Poole, UK). The cells were then washed twice with medium and added to streptavidin coupled magnetic beads at a bead to cell ratio of 30:1. After 30 minutes incubation the beads and rosetted cells were separated by applying the magnetic device three times - taking the supernatant each time.

2.4 DNA/cDNA preparation, PCR amplification and cloning

[0529] DNA was prepared by a simple proteinase-K digest method that was particularly convenient for small numbers of cells (PCR Protocols: A Guide to Methods and Applications. Ed Innis M.A., Gelfand D. H., Sninsky J.J. and White T. J. Academic Press). RNA preparation and subsequent cDNA synthesis was performed as described by Gherardi et al (Gherardi E., Pannell R. and Milstein C. J. *Immunol. Methods*, 1990. 126:61-68). PCR and cloning of the heavy chain libraries was performed using the primers and conditions described by Ward et al (Ward, E.S., Güssow, D., Griffiths, A.D., Jones, P.T. and Winter, G., *Nature*, 1989. 341: 544-546); 40 cycles of PCR amplification were performed. The VH and Fv expression vectors used were adapted from those previously described by Ward et al. They were both subcloned into pUC119 (Veira and Messing see later) and the Fv expression vector was modified to include a germline lambda-1 light chain (obtained as a gift from T. Simon (originally cloned by Siegfried Weiss, Basel Institute of Immunology)). The vector is shown in Figure 53.

2.5 Expression and ELISA

[0530] For screening single colonies were picked into individual wells of microtitre plates (Bibby) in 200µl 2 x TY/ Ampicillin 100µg/ml/0.1% glucose and then incubated at 37°C for 5-6 hours with agitation, Isopropyl-β-D-thiogalactopyranoside (IPTG, Sigma, Poole, UK) was then added to a final concentration of 1 mM and the incubation continued for a further 16 hours at 30°C before harvesting the supernatants. The wells of Falcon ELISA plates (Becton Dickinson, N.J., USA) were coated overnight at room temperature with NIP₁₀-BSA (40µg/ml in PBS) and then blocked with 2% skimmed milk powder in PBS for 2 hours at room temperature. The bacterial supernatants were added and incubated at room temperature for 1 hour and then the plates were washed three times with PBS. Peroxidase conjugated-Goat anti-mouse lambdachain (Southern Biotechnology, Birmingham, USA) was added and again incubated for 1 hour at room temperature before washing six times with PBS and then developing with 2,2'-Azino-bis (3-ethylbenzthiazoline-6-sulfonic acid) (Sigma, Poole, UK) as the peroxidase substrate. The optical density at 405nm was measured using a Thermomax microplate reader (Molecular Devices, Menlo Park, USA) after 30 minutes. Western blotting using the C-terminal myc tag as described in example 23.

3.1 Comparison of RNA/DNA and antigen selected cells

[0531] The results of antigen selection are shown in Table 9. Less than 1% of cells bind to NIP-BSA coated beads and the non-specific binding is very low. Assessment of the proportion of expressed genes from each VH library using western blotting showed that full length VH domains were expressed in 95% (19/20) of all clones when RNA was used as the starting material but only 60% (12/20) of clones when DNA (either selected cells or from total spleen) was used as the starting material. This difference probably results from the fact that many re-arranged pseudogenes could be amplified with our primers and it appears that there must be some degree of selection, at the level of transcription, for functional genes.

[0532] A variable number of clones from each type of library were screened for the production of Fv fragments that bound to NIP. Initial screening ELISAs were performed and positives taken to include those with an optical density of at least twice the background. The initial positives were retransformed and the binding checked in duplicate; it was confirmed that the binding was specific to NIP and not to BSA. The frequency of confirmed positive NIP binding clones for each starting material are shown in Table 10. Using DNA as the starting material for the PCR amplification is approximately equivalent to sampling the cells present as there is only one functional re-arranged heavy chain gene and at most one re-arranged pseudogene per B-cell. Amplifying from the RNA of an animal of course biases the repertoire to the reacting B-cells and in a recently immunised animal this would be expected to give some bias towards the immunogen. The data in Table 10 clearly shows how powerful this selection is with the number of antigen specific genes being enriched at least 96 fold when RNA made one week after primary immunisation is used as the starting material. The data also show that selection for antigen binding cells also provides an alternative powerful method of selection for the required genetic starting material.

3.2 Comparison of Total Spleen/surface immunoglobulin depleted Spleen

[0533] To examine the cellular basis of the selection achieved by using RNA as the starting material we depleted the spleen of surface immunoglobulin positive cells using biotinylated anti-polyvalent immunoglobulin and streptavidin conjugated magnetic beads. Prior FACS analysis had demonstrated that this method removed over 96% of surface immunoglobulin positive cells. RNA was prepared from both surface immunoglobulin depleted and non-depleted fractions of a spleen and VH libraries made from each. The ELISA results (Table 10) show that the number of positives is certainly not decreased by this depletion suggesting that the major portion of the selective effect of using RNA may come from surface immunoglobulin negative G-cells (probably plasma cells).

Conclusions

[0534] The applicants have demonstrated the importance of the amplification of specific RNA produced by immunisation to enable binding activity to be obtained with any reasonable frequency from a combinatorial library. The applicants have also demonstrated an alternative strategy which mimics that of the immune system itself. Using a simple method of selecting for antigen binding cells gave comparable enrichment and has the added advantage of using a broader range of genes. At first sight the random combinatorial approach would appear unlikely to produce the original combination of heavy and light chain because of the vast diversity of the immunoglobulin genes. The applicants show here, however, that following immunisation, with a good antigen, 10% of the VH genes from total splenic RNA isolated come from antigen specific cells so the effective size of the repertoire is greatly reduced. This together with the fact that promiscuity of the heavy and light chains occurs (examples 17 and 18) accounts for the fact that combinatorial system does produce antigen binding clones with reasonable frequency. The data also suggests that the bulk of the antigen specific RNA comes from surface immunoglobulin negative cells which are most likely plasma cells.

[0535] The data also show that this simple method of antigen selection may be useful in reducing the complexity of the combinatorial library. In this case an enrichment of antigen specific genes of at least 56 fold has been achieved which in the normal case where heavy and light chains are unknown would result in a reduction of the complexity of the combinatorial library by a factor of over 3000. A further advantage of using antigen selected cells (and amplifying from DNA to reduce any bias due to the state of the cell) is that this results in a broader range of antibody genes amplified. It may be that a simple cell selection such as that the applicants have described here in combination with phage selection would be ideal. From this example it can be seen that by combining cell and phage selection methods one could reasonably expect to screen all the combinations of heavy and light chain (approximately 4×10^{10}) and would thus be able to screen all binding combinations although this would not, at present, be possible from whole spleen (approximately 4×10^{14} combinations, assuming 50% B-cells).

Table 1.

Enrichment of pAb (D1.3) from vector population			
INPUT RATIO ^a	OUTPUT RATIO		ENRICHMENT ^d
	oligo ^b	ELISA ^c	
pAb:fd-CAT1	pAb:total phage	pAb:total phage	
Single Round			
1:4x10 ³	43/124		1.3x10 ³
1:4x10 ⁴	2/82		1.0x10 ³
Two Rounds			
1:4x10 ⁴	197/372		2.1x10 ⁴
1:4x10 ⁵	90/356	3/24	1.0x10 ⁵
1:4x10 ⁶	27/183	5/26	5.9x10 ⁵
1:4x10 ⁷	13/278		1.8x10 ⁶

Footnotes: ^aApproximately 10¹² phage with the stated ratio of pAb (D1.3) : FDTPs/Bs were applied to 1 ml lysozyme-sepharose columns, washed and eluted.

^bTG1 cells were infected with the eluted specific binding phage and plated onto TY-tet plates. After overnight incubation at 30-37°C, the plates were analysed by hybridisation to the ³²p, labelled oligonucleotide VH1FOR (Ward et al op cit) which is specific to pAb D1.3.

^cSingle colonies from overnight plates were grown, phage purified, and tested for lysozyme binding.

^dEnrichment was calculated from the oligonucleotide probing data.

Table 2

Enrichment of pAb (D1.3) from mixed pAb population		
Input Ratio ¹ (pAbD1.3:pAbNQ11)	Output Ratio ² (pAb D1.3:Total phage)	Enrichment
Single Round		
1 : 2.5 x 10 ⁴	18/460	0.98 x 10 ³
1 : 2.5 x 10 ⁵	3/770	- 0.97 x 10 ³
1 : 2.5 x 10 ⁶	0/112	-
pAb NQ11 only	0/460	-
Second Round		
1 : 2.5 x 10 ⁴	119/170	1.75 x 10 ⁴
1 : 2.5 x 10 ⁵	101/130	1.95 x 10 ⁵
1 : 2.5 x 10 ⁶	102/204	1.26 x 10 ⁶
1 : 2.5 x 10 ⁷	0/274	-
1 : 2.5 x 10 ⁸	0/209	-
pAb NQ11 only	0/170	-

Notes

1. 10¹⁰ phage applied to a lysozyme column as in table 1.

2. Plating of cells and probing with oligonucleotide as in table 1, except the oligonucleotide was D1.3CDR3A.

Table 3. Affinity selection of hapten-binding phage.

		Clones binding to phOx [†]		
third round		Pre-column	After first round	After second round
A Random Combinatorial Libraries				
	phOx-immunised mice	0/568 (0%)	48/376 (13%)	175/188 (93%)
	Unimmunised mice			0/388 (0%)
B Hierarchical Libraries				
	VH-B/Vκ- <i>rep</i> library	6/190 (3%)	348/380 (92%)	
	VH- <i>rep</i> /Vκ-d library	0/190 (0%)	23/380 (7%)	
C Fractionation of VH-B/Vκ-d and VH-B/Vκ-b phage[†]				
	Mixture of clones	88/1896 (4.6%) [44/1740 (2.5%)*]	55/95 (57.9%)	1152/1156 (99.7%)
				1296/1299 (99.8%)

* In panel C, numbers refer to VH-B/Vκ-d colonies.

† Numbers after three reinfections and cycles of growth. This control, omitting the column steps, confirms that a spurious growth or infectivity advantage was not responsible for the enrichment for clone VH-B/Vκ-d

Table 4

Phage/Phagemid [†]	Helper Phage	Binding to phOx [*]	Chain(s) displayed [#]	Chain as gene III fusion [*]	Soluble chain(s) [#]
A					
fd CAT2		non binding	none		
fd CAT2-I		binding	scFv	scFv	heavy chain
fd CAT2-II		binding	Fab	light chain	
pHEN1	VCSMB	non binding	none		
pHEN1 I	VCSMB	binding	scFv	scFv	heavy chain
pHEN1 II	VCSMB	binding	Fab	light chain	
B					
pHEN1 I (HB2151)		binding			scFv [§]
pHEN1 II (HB2151)		binding			Fab [§]
C					
fd CAT2-III		non binding	heavy chain	heavy chain	heavy chain
fd CAT2-IV		non binding	light chain	light chain	heavy chain
pHEN1 III (HB2151)	VCSMB	non binding	none		heavy chain
pHEN1 III (HB2151)	fd-tet-DOG1 IV	binding	Fab	light chain	heavy chain
pHEN1 IV (HB2151)	VCSMB	non binding	none		light chain
pHEN1 IV (HB2151)	fd-tet-DOG1 III	binding	Fab	heavy chain	light chain

Overview of phOx-BSA ELISA results of phage and phagemid constructions.

^{*} Phase were considered to be 'binding' if OD₄₀₅ of sample was at least 10 fold greater than background in ELISA; [†] *E. coli* TG1 was used for the growth of the phage unless the use of *E. coli* HB2151 is specifically indicated; [#] Information deduced from genetic structure and in accordance with binding data; [§] Result confirmed experimentally by Western blot (for Fab, see

Figure 26.

Table 5

Mutations in XscFtB18 selected by display on phage following growth in mutator strains		
Nucleotide mutation (base position)	Amino acid mutation	Number
308	Ala->Val (VH FR3)	3

EP 0 774 511 B1

Table 5 (continued)

Mutations in XscFtB18 detected by display on phage following growth in mutant strains		
Nucleotide mutation (base position)	Amino acid mutation	Number
703	Tyr->Asp (VL CDR3)	1
706	Ser-> Gly (VL CDR3)	1
724	Gly-> Ser (VL FR4)	21
725	Gly-> Asp (VL FR4)	3
734	Thr-> Ile (VL FR4)	1

Table 6(i) Oligonucleotide primers used for PCR of human immunoglobulin genes

Oligo Name	Sequence
Human VH Back Primers	
HuVH1aBACK	5'-CAG GTG CAG CTG GTG CAG TCT GG-3'
HuVH2aBACK	5'-CAG GTC AAC TTA AGG GAG TCT GG-3'
HuVH3aBACK	5'-GAG GTG CAG CTG GTG GAG TCT GG-3'
HuVH4aBACK	5'-CAG GTG CAG CTG CAG GAG TCG GG-3'
HuVH5aBACK	5'-GAG GTG CAG CTG TTG CAG TCT GC-3'
HuVH6aBACK	5'-CAG GTA CAG CTG CAG CAG TCA GG-3'
HuVH1aBACKSfi	5'-GTC CTC GCA ACT GCG GCC CAG CCG GCC ATG GCC CAG GTG CAG CTG GTG CAG TCT GG-3'
HuVH2aBACKSfi	5'-GTC CTC GCA ACT GCG GCC CAG CCG GCC ATG GCC CAG GTC AAC TTA AGG GAG TCT GG-3'
HuVH3aBACKSfi	5'-GTC CTC GCA ACT GCG GCC CAG CCG GCC ATG GCC GAG CTG GTG GAG TCT GG-3'
HuVH4aBACKSfi	5'-GTC CTC GCA ACT GCG GCC CAG CCG GCC ATG GCC CAG CTG CAG GAG TCG GG-3'
HuVH5aBACKSfi	5'-GTC CTC GCA ACT GCG GCC CAG CCG GCC ATG GCC CAG CTG TTG CAG TCT GC-3'
HuVH6aBACKSfi	5'-GTC CTC GCA ACT GCG GCC CAG CCG GCC ATG GCC CAG GTA CAG CTG CAG TCA GG-3'
Human JH Forward Primers	
HuJH1-2FOR	5'-TGA GGA GAC GGT GAC CAG GGT GCC-3'
HuJH3FOR	5'-TGA AGA GAC GGT GAC CAT TGT CCC-3'
HuJH4-5FOR	5'-TGA GGA GAC GGT GAC CAG GGT TCC-3'
HuJH6FOR	5'-TGA GGA GAC GGT GAC CGT GGT CCC-3'
Human Heavy Chain Constant Region Primers	

Table 6(ii)

HuIgG1-4CH1FOR 5'-GTC CAC CTT GGT GTT GCT GGG CTT-3'
 HuIgMFOR 5'-TGG AAG AGG CAC GTT CTT TTC TTT-3'

Human V κ Back Primers

HuV κ 1aBACK 5'-GAC ATC CAG ATG ACC CAG TCT CC-3'
 HuV κ 2aBACK 5'-GAT GTT GTG ATG ACT CAG TCT CC-3'
 HuV κ 3aBACK 5'-GAA ATT GTG TTG ACG CAG TCT CC-3'
 HuV κ 4aBACK 5'-GAC ATC GTG ATG ACC CAG TCT CC-3'
 HuV κ 5aBACK 5'-GAA ACG ACA CTC ACG CAG TCT CC-3'
 HuV κ 6aBACK 5'-GAA ATT GTG CTG ACT CAG TCT CC-3'

Human J κ Forward Primers

HuJ κ 1FOR 5'-ACG TTT GAT TTC CAC CTT GGT CCC-3'
 HuJ κ 2FOR 5'-ACG TTT GAT CTC CAG CTT GGT CCC-3'
 HuJ κ 3FOR 5'-ACG TTT GAT ATC CAC TTT GGT CCC-3'
 HuJ κ 4FOR 5'-ACG TTT GAT CTC CAC CTT GGT CCC-3'
 HuJ κ 5FOR 5'-ACG TTT AAT CTC CAG TCG TGT CCC-3'
 HuJ κ 1BACKNot 5'-GAG TCA TTC TCG ACT TGC GGC CGC ACG TTT GAT TTC CAC CTT GGT CCC-3'
 HuJ κ 2BACKNot 5'-GAG TCA TTC TCG ACT TGC GGC CGC ACG TTT GAT CTC CAG CTT GGT CCC-3'
 HuJ κ 3BACKNot 5'-GAG TCA TTC TCG ACT TGC GGC CGC ACG TTT GAT ATC CAC TTT GGT CCC-3'
 HuJ κ 4BACKNot 5'-GAG TCA TTC TCG ACT TGC GGC CGC ACG TTT GAT CTC CAC CTT GGT CCC-3'
 HuJ κ 5BACKNot 5'-GAG TCA TTC TCG ACT TGC GGC CGC ACG TTT AAT CTC CAG TCG TGT CCC-3'

Human κ Constant Region Primers

Table 6(iii)

HuCrFOR	5'-AGA CTC TCC CCT GTT GAA GCT CTT-3'
HuCrFORNot1	5'-GAG TCA TTC TCG ACT TGC GGC CGC TTA TTA AGA CTC TCC CCT GTT GAA GCT CTT-3'
HuCrFORNot2	5'-GAG TCA TTC TCG ACT TGC GGC CGC AGA CTC TCC CCT GTT GAA GCT CTT-3'

Human λ Back Primers

HuV λ 1BACK	5'-CAG TCT GTG TTG ACG CAG CCG CC-3'
HuV λ 2BACK	5'-CAG TCT GCC CTG ACT CAG CCT GC-3'
HuV λ 3aBACK	5'-TCC TAT GTG CTG ACT CAG CCA CC-3'
HuV λ 3bBACK	5'-TCT TCT GAG CTG ACT CAG GAC CC-3'
HuV λ 4BACK	5'-CAC GTT ATA CTG ACT CAA CCG CC-3'
HuV λ 5BACK	5'-CAG GCT GTG CTC ACT CAG CCG TC-3'
HuV λ 6BACK	5'-AAT TTT ATG CTG ACT CAG CCC CA-3'

Human λ Forward Primers

HuJ λ 1FOR	5'-ACC TAG GAC GGT GAC CTT GGT CCC-3'
HuJ λ 2-3FOR	5'-ACC TAG GAC GGT CAG CTT GGT CCC-3'
HuJ λ 4-5FOR	5'-ACC TAA AAC GGT GAG CTG GGT CCC-3'

HuJ λ FORNOT	5'-GAG TCA TTC TCG ACT TGC GGC CGC ACC TAG GAC GGT GAC CTT GGT CCC-3'
HuJ λ 2-3FORNOT	5'-GAG TCA TTC TCG ACT TGC GGC CGC ACC TAG GAC GGT CAG CTT GGT CCC-3'
HuJ λ 4-5FORNOT	5'-GAG TCA TTC TCG ACT TGC GGC CGC ACY TAA AAC GGT GAG CTG GGT CCC-3'

Human λ Constant Region Primers

Table 6(iv)

Hu λ FOR	5'-TGA AGA TTC TGT AGG GGC CAC TGT CTT-3'
Hu λ FORNot1	5'-GAG TCA TTC TCG ACT TGC GGC CGC TTA TTA TGA AGA TTC TGT AGG GGC CAC TGT CTT-3'
Hu λ FORNot2	5'-GAG TCA TTC TCG ACT TGC GGC CGC TGC AGA TTC TGT AGG GGC TGT CTT-3'

Linker oligos

Reverse JH for scFv linker

RHuJH1-2	5'-GCA CCC TGG TCA CCG TCT CCT CAG GTG G-3'
RHuJH3	5'-GGA CAA TGG TCA CCG TCT CTT CAG GTG G-3'
RHuJH4-5	5'-GAA CCC TGG TCA CCG TCT CCT CAG GTG G-3'
RHuJH6	5'-GGA CCA CCG TCA CCG TCT CCT CAG GTG C-3'

Reverse IgG1-4CH1 primer for Fab linker

RHuIgG1-4CH1FOR	5'-AAG CCC AGC AAC ACC AAG GTG GAC-3'
-----------------	---------------------------------------

Reverse V κ for scFv linker

RHuV κ 1aBACKFv	5'-GGA GAC TGG GTC ATC TGG ATG TCC GAT CCG CC-3'
RHuV κ 2aBACKFv	5'-GGA GAC TGA GTC ATC ACA ACA TCC GAT CCG CC-3'
RHuV κ 3aBACKFv	5'-GGA GAC TGC GTC AAC ACA ATT TCC GAT CCG CC-3'
RHuV κ 4aBACKFv	5'-GGA GAC TGG GTC ATC ACG ATG TCC GAT CCG CC-3'
RHuV κ 5aBACKFv	5'-GGA GAC TGC GTG AGT GTC GTT TCC GAT CCG CC-3'
RHuV κ 6aBACKFv	5'-GGA GAC TGA GTC AGC ACA ATT TCC GAT CCG CC-3'

Reverse V κ for Fab linker

Table 6(v)

RHuVκ1aBACKFab	5'-GGA GAC TGG GTC ATC TGG ATG TCG GCC ATC GCT GG-3'
RHuVκ2aBACKFab	5'-GGA GAC TGC GTC ATC ACA ACA TCG GCC ATC GCT GG-3'
RHuVκ3aBACKFab	5'-GGA GAC TGC GTC AAC ACA ATT TCG GCC ATC GCT GG-3'
RHuVκ4aBACKFab	5'-GGA GAC TGG GTC ATC ACG ATG TCG GCC ATC GCT GG-3'
RHuVκ5aBACKFab	5'-GGA GAC TGC GTC AGT GTC GTT TCG GCC ATC GCT GG-3'
RHuVκ6aBACKFab	5'-GGA GAC TGC GTC AGC ACA ATT TCG GCC ATC GCT GG-3'

Reverse Vλ for svFv linker

RHuVλBACK1Fv	5'-GGC GGC TGC GTC AAC ACA GAC TGC GAT CCG CCA CCG CCA GAG-3'
RHuVλBACK2Fv	5'-GCA GGC TGA GTC AGA GCA GAC TGC GAT CCG CCA CCG CCA GAG-3'
RHuVλBACK3aFv	5'-GGT GGC TGA GTC AGC ACA TAG GAC GAT CCG CCA CCG CCA GAG-3'
RHuVλBACK3bFv	5'-GGG TCC TGA GTC AGC TCA GAA GAC GAT CCG CCA CCG CCA GAG-3'
RHuVλBACK4Fv	5'-GGC GGT TGA GTC AGT ATA ACG TGC GAT CCG CCA CCG CCA GAG-3'
RHuVλBACK5Fv	5'-GAC GGC TGA GTC AGC ACA GAC TGC GAT CCG CCA CCG CCA GAG-3'
RHuVλBACK6Fv	5'-TGG GGC TGA GTC AGC ATA AAA TTC GAT CCG CCA CCG CCA GAG-3'

Reverse Vλ for Fab linker

RHuVλBACK1Fab	5'-GGC GGC TGC GTC AAC ACA GAC TGG GCC ATC GCT GGT TGG GCA-3'
RHuVλBACK2Fab	5'-GCA GGC TGA GTC AGA GCA GAC TGG GCC ATC GCT GGT TGG GCA-3'
RHuVλBACK3aFab	5'-GGT GGC TGA GTC AGC ACA TAG GAG GCC ATC GCT GGT TGG GCA-3'
RHuVλBACK3bFab	5'-GGG TCC TGA GTC AGC TCA GAA GAG GCC ATC GCT GGT TGG GCA-3'
RHuVλBACK4Fab	5'-GGC GGT TGA GTC AGT ATA ACG TGG GCC ATC GCT GGT TGG GCA-3'
RHuVλBACK5Fab	5'-GAC GGC TGA GTC AGC ACA GAC TGG GCC ATC GCT GGT TGG GCA-3'
RHuVλBACK6Fab	5'-TGG GGC TGA GTC AGC ATA AAA TTG GCC ATC GCT GGT TGG GCA-3'

Table 7 Deduced protein sequences of heavy and light chains selected from unimmunized library

Oxazolone binder

HEAVY CHAIN
 VH15.4 QVQLVQSGAEVKKPGASVKVSCKASQYPTT SYGLS WVRQAPGQGLEWV HIGAVKENTKYAQKLQD RVTMTITVISTVYHMLASLASDITAVYYCVR LIPRTATLAI YVINDVAKKEI

LIGHT CHAIN
 VL15.4 NAYYS WYQLPCTAPRLIY DNRKPS QIPDRFSQSKSTISNTLITLGLQVNDIAYYC GTWDER

BSA binders

HEAVY CHAIN
 VH3.5 QVQLVQSGDGVVQPRSLSLSCASQFTTS SYGMH WVRQAPGQGLEWV VISYDGSNTYNDVKG RFTTISRDNSNITLVQVHSLRABDTAVYYCAR TQYSSGRDY FDMHCOOT

LIGHT CHAIN
 VL3.5 SSELTPDPANVSVALQOTVRIIC QGDSLRYYAS WYQQKPGQAPFLVIY QGNRPIS QIPDRFGSSSSNTLITITDQAQDEADYYC HSRDSSGIM WVPQD

Lysuzyme binders:

HEAVY CHAINS

VH10.1 ELTCSVBCDSIS EOOTYR WIRQSPCKGLEHIG SYHSDPTTYNPSLKS RVTNSVDTSKQFSLKLSVTAADTAVYYCAR EGGSTMRSILYMI YMDVHOK
 VH14.1 QVQLVQSGAEVKKPSETLSLVCYBOGSL FBYMG WIRQPPCKGLEHIG VISIRKTDYNGSLQS RVTIGADTSKQFSLKLSVTAADTAVYYCAR SFNSIFFFOY HCOOT
 VH13.1 QVQLVQSGAEVKKPQSLHISQCSYGFIS NYMIG WVRQPPCKGLEHIG ILYPQSDTRYSPSFQD QVTIGADKISITAVYIIMSSLSASDTALYYCAR LVCOOTPAY HCOOT
 VH16.1 QVQLVQSGAEVKKPQSLRISCKDGYEFS TYNIG WVRQPPCKGLEHIG ILYPQSDTRYSPSFQD QVTISVDISITITAYIIMSSLSA

LIGHT CHAINS

VH10.1 EIVLTQSPSSLSNVGDRTVITC RASQSIHNTLV WYQQKPGQAPRLIY NASTLOS QVPSRESGCSGTFTLTINSIQPDEPATYYC QQTIIFFP LTFQGG
 VH14.1 SSELTPDPANVSVALQOTVRIIC QGDSLRBSYAS WYQQKPGQAPFLVIY QENRPIS QIPDRFGSSSKVTSLSLITDQCRDEADYYC NSRDSRCTAL EVPQD
 VH13.1 HVILTOPASVSGSPQSITIEC TQSSRDVCOITNVIS WYQHHPQAPRLIIS EVNRPIS QVNSRESGSKSVTSLSLITISLQCRDEADYYC NSYTSSEKT YVPQD
 VH16.1 QSNLTQPAVSGSPQSITIEC SGSSSDIQRDYNVS WYQINPDAPRLIY EVNRPIS QISIRFENSKECHTSLTIGELQCRDEADYYC NSYT

Table 8

Enrichment of pAbNQ11 from pAbD1.3 background by affinity selection using Ox-BSA biotinylated with a cleavable reagent and binding to streptavidin magnetic beads		
Input Ratio ¹ (pAbD1.3:pAbNQ11)	Output Ratio ² (pAb NQ11: Total phage)	Enrichment
2235:1	61/197	690
22350:1	5/202	544

1. 1.9×10^{10} phage in 0.5ml mixed for 1 hour with 5 μ l streptavidin-magnetic heads precoated with antigen (OX-BSA).

2. Colonies probed with the oligonucleotide NQ11CDR3

Table 9:

Results of antigenic cell selection		
	Number of Cells	% of total cells
Total spleen cells	4x10 ⁷ -	
Cells bound to uncoated beads	0.8x10 ⁴	0.02
Cells bound to NIP-BSA coated beads	22x10 ⁴	0.55

Table 10:

Results of Fv MP binding ELISAs from selected cell populations:		
	Positives	*Degree of Enrichment
Cell Population DNA from total spleen	0/940	-
RNA from total Spleen	29/282	> 96
DNA from antigen binding cells	17/282	>56
Surface Ig Selection		
RNA from Surface Ig negative fraction	8/94	-
RNA from total Spleen	4/94	-

*Degree of enrichment compared to total DNA.

Claims

1. A filamentous bacteriophage particle displaying on its surface a binding molecule which is a Fab or single-chain Fv antibody molecule comprising a binding domain able to bind target epitope or antigen, the particle containing a phagemid genome comprising nucleic acid with a nucleotide sequence encoding the binding molecule expressed from the nucleic acid and displayed by the particle at its surface.
2. A filamentous bacteriophage particle according to claim 1 wherein said binding molecule is a scFv antibody molecule.
3. A filamentous bacteriophage particle according to claim 1 wherein said binding molecule is a Fab antibody molecule.
4. A population of filamentous bacteriophage particles according to any one of claims 1 to 3 displaying a population of said binding molecules having a range of binding specificities.

5. A method for producing a filamentous bacteriophage particle displaying at its surface a binding molecule specific for a particular target epitope or antigen, which method comprises the steps of:

producing a population of filamentous bacteriophage particles displaying at their surface a population of binding molecules having a range of binding specificities, wherein the binding molecules are Fab antibody molecules able to bind target epitope or antigen, and wherein each filamentous bacteriophage particle contains a phagemid genome comprising nucleic acid with a nucleotide sequence encoding the binding molecule expressed from the nucleic acid and displayed by the particle at its surface;

selecting for a filamentous bacteriophage particle displaying a binding molecule with a desired specificity by contacting the population of filamentous bacteriophage particles with a target epitope or antigen so that individual binding molecules displayed on filamentous bacteriophage particles with the desired specificity bind to said target epitope or antigen.

6. A method according to claim 5 additionally comprising separating bound filamentous bacteriophage particles from the target epitope or antigen.

7. A method according to claim 6 additionally comprising recovering separated filamentous bacteriophage particles displaying a binding molecule with the desired specificity.

8. A method for producing a binding molecule specific for a particular target epitope or antigen, which method comprises:

performing the method according to claim 7;
isolating from separated filamentous bacteriophage particles recovered according to the method of claim 7 nucleic acid encoding the binding molecule;
inserting nucleic acid encoding the binding molecule, or a fragment or derivative thereof with binding specificity for the target epitope or antigen, in a recombinant system; and
producing the binding molecule, or fragment or derivative thereof with binding specificity for the target epitope or antigen, in the recombinant system separate from filamentous bacteriophage particles.

Patentansprüche

1. Filamentöses Bakteriophagen-Teilchen, auf dessen Oberfläche ein Bindungsmolekül präsentiert wird, das ein Fab- oder Einzelketten-Fv-Antikörpermolekül ist, das eine Bindungsdomäne umfasst, die sich an Target-Epitop oder -Antigen binden kann, wobei das Teilchen ein Phagemid-Genom enthält, das Nucleinsäure mit einer Nucleotidsequenz umfasst, die für das Bindungsmolekül kodiert, das von der Nucleinsäure exprimiert wird und auf der Oberfläche des Teilchens präsentiert wird.
2. Filamentöses Bakteriophagen-Teilchen nach Anspruch 1, worin das Bindungsmolekül ein scFv-Antikörpermolekül ist.
3. Filamentöses Bakteriophagen-Teilchen nach Anspruch 1, worin das Bindungsmolekül ein Fab-Antikörpermolekül ist.
4. Population von filamentösen Bakteriophagen-Teilchen nach einem der Ansprüche 1 bis 3, auf denen eine Population der Bindungsmoleküle präsentiert wird, die eine Bandbreite von Bindungsspezifitäten aufweisen.
5. Verfahren zur Herstellung eines filamentösen Bakteriophagen-Teilchens, auf dessen Oberfläche ein Bindungsmolekül präsentiert wird, das für ein bestimmtes Target-Epitop oder -Antigen spezifisch ist, wobei das Verfahren folgende Schritte umfasst:

das Erzeugen einer Population filamentöser Bakteriophagen-Teilchen, auf deren Oberfläche eine Population von Bindungsmolekülen präsentiert wird, die eine Bandbreite von Bindungsspezifitäten aufweisen, worin die Bindungsmoleküle Fab-Antikörpermoleküle sind, die sich an das Target-Epitop oder -Antigen binden können, und worin jedes filamentöse Bakteriophagen-Teilchen ein Phagemid-Genom enthält, das Nucleinsäure mit einer Nucleotidsequenz umfasst, die für das Bindungsmolekül kodiert, das von der Nucleinsäure exprimiert

wird und auf der Oberfläche der Teilchen präsentiert wird,
das Selektieren bezüglich eines filamentösen Bakteriophagen-Teilchens, auf dem ein Bindungsmolekül mit
einer gewünschten Spezifität präsentiert wird, durch Kontaktieren der Population filamentöser Bakteriophagen-Teilchen mit einem Target-Epitop oder -Antigen, so dass einzelne Bindungsmoleküle, die auf den filamentösen Bakteriophagen-Teilchen präsentiert werden, sich mit der gewünschten Spezifität an das Target-Epitop oder -Antigen binden.

6. Verfahren nach Anspruch 5, das zusätzlich das Abtrennen gebundener filamentöser Bakteriophagen-Teilchen vom Target-Epitop oder -Antigen umfasst.

7. Verfahren nach Anspruch 6, das zusätzlich das Gewinnen abgetrennter filamentöser Bakteriophagen-Teilchen umfasst, auf denen ein Bindungsmolekül mit der gewünschten Spezifität präsentiert wird.

8. Verfahren zur Herstellung eines Bindungsmoleküls, das für ein bestimmtes Target-Epitop oder -Antigen spezifisch ist, wobei das Verfahren umfasst:

die Durchführung des Verfahrens nach Anspruch 7;
das Isolieren von Nucleinsäure, die für das Bindungsmolekül kodiert, von abgetrennten filamentösen Bakteriophagen-Teilchen, die nach dem Verfahren nach Anspruch 7 gewonnen wurden;
das Insertieren von Nucleinsäure, die für das Bindungsmolekül kodiert, oder eines Fragments oder Derivats davon mit Bindungsspezifität für das Target-Epitop oder -Antigen, in einem rekombinanten System; und
das Erzeugen des Bindungsmoleküls oder Fragments oder Derivats davon mit Bindungsspezifität für das Target-Epitop oder -Antigen im rekombinanten System, getrennt von den filamentösen Bakteriophagen-Teilchen.

Revendications

1. Particule bactériophage filamentaire présentant à sa surface une molécule de liaison qui est une molécule d'anticorps Fab ou Fv à une seule chaîne comprenant un domaine de liaison capable de lier un épitope cible ou antigène, la particule contenant un génome phagémide comprenant de l'acide nucléique avec une séquence de nucléotides codant pour la molécule de liaison exprimée à partir de l'acide nucléique et présentée par la particule à sa surface.

2. Particule -bactériophage filamentaire selon la revendication 1 où ladite molécule de liaison est une molécule d'anticorps scFv.

3. Particule bactériophage filamentaire selon la revendication 1 où ladite molécule de liaison est une molécule d'anticorps Fab.

4. Population de particules bactériophages filamentaires selon l'une quelconque des revendications 1 à 3 présentant une population desdites molécules de liaison ayant une gamme de spécificités de liaison.

5. Méthode de production d'une particule bactériophage filamentaire présentant à sa surface une molécule de liaison spécifique d'un épitope cible ou antigène particulier, laquelle méthode comprend les étapes de :

produire une population de particules bactériophages filamentaires présentant à leur surface une population de molécules de liaison ayant une gamme de spécificités de liaison, où les molécules de liaison sont des molécules d'anticorps Fab capables de lier l'épitope cible ou antigène, et où chaque particule bactériophage filamentaire contient un génome phagémide comprenant de l'acide nucléique avec une séquence de nucléotides codant pour la molécule de liaison exprimée par l'acide nucléique et présentée par la particule à sa surface;

sélectionner pour une particule bactériophage filamentaire présentant une molécule de liaison avec une spécificité souhaitée par mise en contact de la population des particules bactériophages filamentaires avec un épitope cible ou antigène de façon que des molécules individuelles de liaison présentées sur les particules bactériophages filamentaires avec la spécificité souhaitée se lient audit épitope cible ou antigène.

6. Méthode selon la revendication 5 comprenant additionally

la séparation des particules bactériophages filamenteux liées de l'épitope cible ou antigène.

7. Méthode selon la revendication 6 comprenant additionally

la récupération des particules bactériophages filamenteux séparées présentant une molécule de liaison avec la spécificité souhaitée.

8. Méthode de production d'une molécule de liaison spécifique d'un épitope cible ou antigène particulier, laquelle méthode comprend:

l'accomplissement de la méthode selon la revendication 7;

l'isolement de particules bactériophages filamenteux séparées, récupérées selon la méthode de la revendication 7 de l'acide nucléique codant pour la molécule de liaison; et

l'insertion de l'acide nucléique codant pour la molécule de liaison ou d'un fragment ou dérivé de celui-ci avec une spécificité de liaison pour l'épitope cible ou antigène, dans un système recombinant; et

la production de la molécule de liaison, ou fragment ou dérivé de celle-ci avec une spécificité de liaison pour l'épitope cible ou antigène, dans le système recombinant séparé des particules bactériophages filamenteux.

Fig. 1.

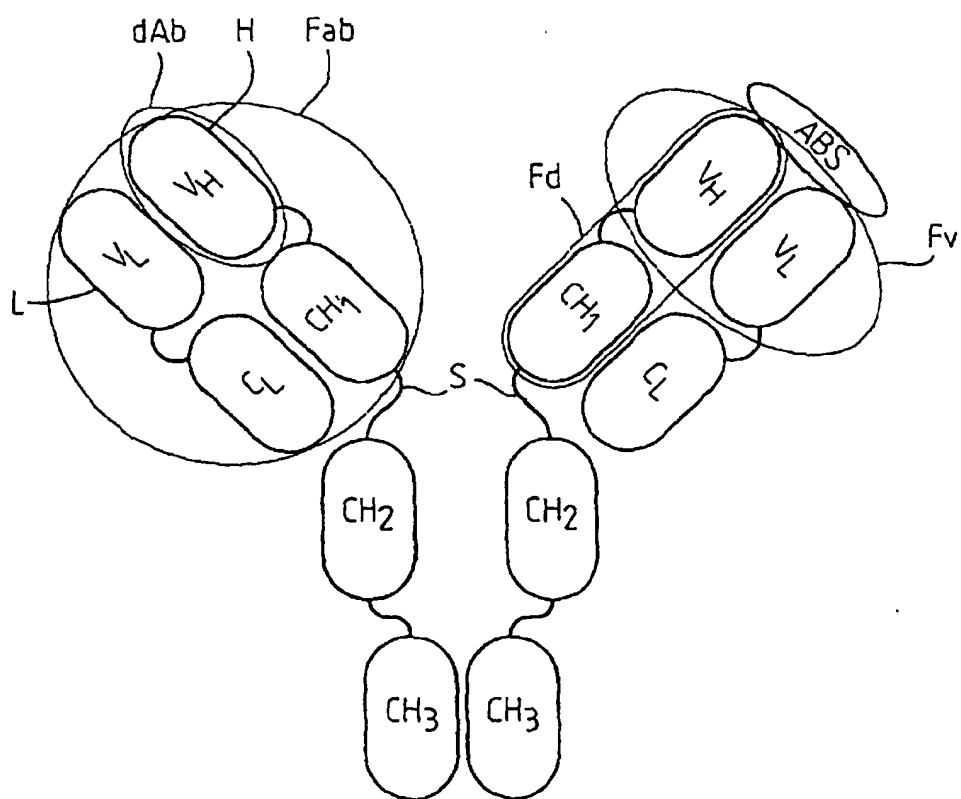


Fig. 2 (i)

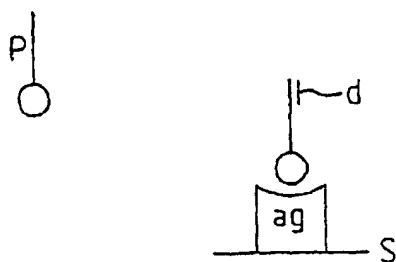
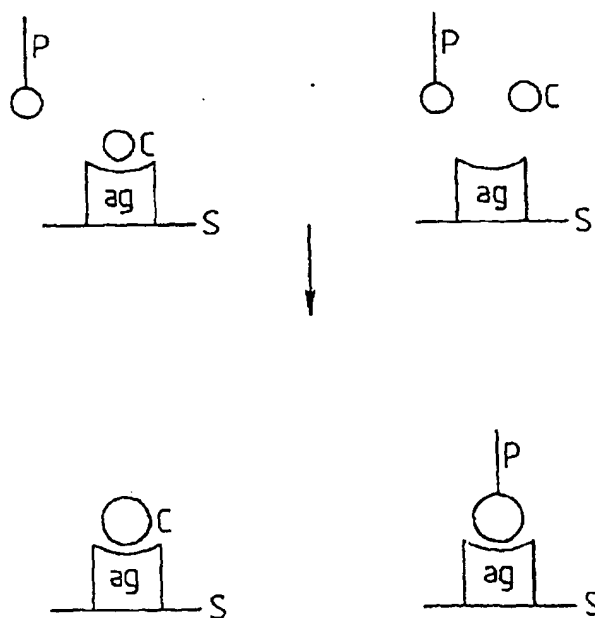
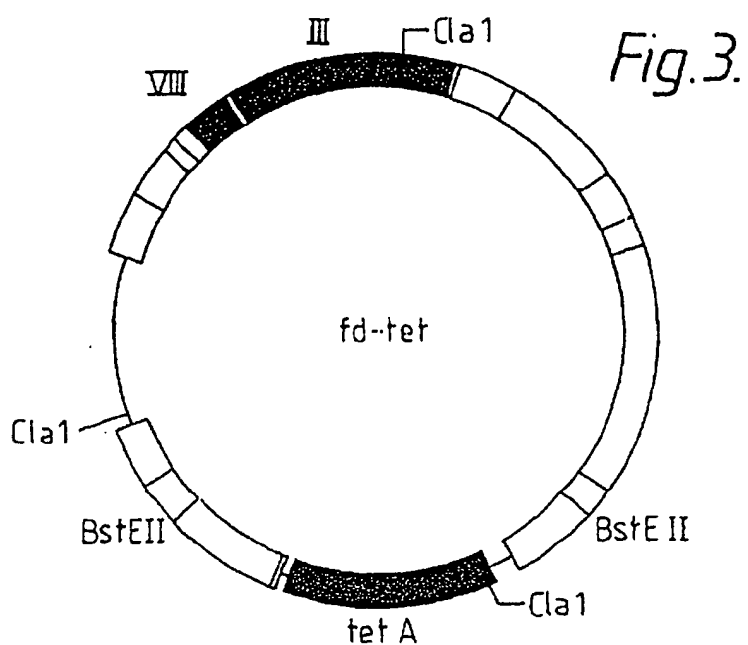


Fig. 2 (ii)





fd - tet
 ~
 cleave with BstEII
 ~
 fill in with Klenow
 ~
 re-ligate
 ↓
 FDTδBst
 ~
 in vitro mutagenesis (oligo 1)
 ↓
 FDTPs/Bs
 ~
 in vitro mutagenesis (oligo 2)
 ↓
 FDTPs/Xh

Fig. 4.1

Oligo 1 (1653) ACA ACT TTC AAC AGT TGA GGA GAC GGT GAC CGT AAG CTT CTG CAG TTG GAC CTG AGC
GGA GTG AGA ATA (1620)

Oligo 2 (1653) ACA ACT TTC AAC AGT TTC CCG TTT GAT CTC GAG CTC CTG CAG TTG GAC CTG

Oligo 3 (1704) GTC GTC TTT CCA GAC GTT AGT

Fig. 4.2

GENE III

GENE III

(1624) A TCT CAC TCC GCT

(1650) GAA ACTGTT GAA AGT

SIGNAL
CLEAVAGE SITE

Q V Q L Q V T V S S

B TCT CAC TCC GCT CAG GTC CAA CTG CAG AAG CTT ACG GTC ACC GTC TCC TCA ACT GTT GAA AGT
PstI BstEII

Q V Q L Q L E I K R

C TCT CAC TCC GCT CAG GTC CAA CTG CAG GAG CTC GAG ATC AAA CCG GAA ACT GTT GAA AGT
PstI XhoI

Fig. 5.

rbs M K Y L L P T A A
 GCATGCAAAATTCCTATTTCAAGGAGACAGTCATAATGAAATACCTATTGCCTACGGCAGCC
 10 20 30 40 50 60
 SphI
 PelB leader
 A G L L L L A A O P A M A Q V Q L Q E S
 GCTGGATTGTTATTACTCGCTGCCCAACCAGCGATGGCCCAAGTGCAGCTGCAGGAGTCA
 70 80 90 100 110 120
 PstI
 G P G L V A P S Q S L S I T C T V S G F
 GGACCTGGCCTGGTGGCGCCCTCACAGAGCCTGTCCATCACATGCACCGTCTCAGGGTTC
 130 140 150 160 170 180
 S L T G Y G V N W V R Q P P G K G L E W
 TCATTAACCGGCTATGGTGTAAACTGGGTTCGCCAGCCTCCAGGAAAGCGTCTGGAGTGG
 190 200 210 220 230 240
 VHD1.3
 L G M I W G D G N T D Y N S A L K S R L
 CTGGGAATGATTTGGGGTGATGGAAACACAGACTATAATTCAGCTCTCAAATCCAGACTG
 250 260 270 280 290 300
 S I S K D N S K S Q V F L K M N S L H T
 AGCATCAGCAAGGACAACCTCCAAGAGCCAAGTTTTCTTAAAAATGAACAGTCTGCACACT
 310 320 330 340 350 360
 D D T A R Y Y C A R E R D Y R L D Y W G
 GATGACACAGCCAGGTACTACTGTGCCAGAGAGAGAGATTATAGGCTTGACTACTGGGGC
 370 380 390 400 410 420
 Linker Peptide
 Q G T T V T V S S G G G G S G G G G S G
 CAAGGCACCACGGTCACCGTCTCCTCAGgtggaggcgggttcaggcggaggtgggtctggc
 430 440 450 460 470 480
 HstEII
 G G G S D I E L T Q S P A S L S A S V G
 ggtggcggatcgGACATCGAGCTCACTCAGTCTCCAGCCTCCCTTTCTGCGTCTGTGGGA
 490 500 510 520 530 540
 SacI

Fig. 5 cont.

E T V T I T C R A S G N I H N Y L A W Y
 GAAACTGTCACCATCACATGTCGAGCAAGTGGGAATATTCACAATTATTTAGCATGGTAT
 550 560 570 580 590 600

Q Q K Q G K S P Q L L V Y Y T T T L A D
 CAGCAGAAACAGGGAAAATCTCCTCAGCTCCTGGTCTATTATACAACAACCTTAGCAGAT
 610 620 630 640 650 660

VKD1.3

G V P S R F S G S G S G T Q Y S L K I N
 GGTGTGCCATCAAGGTTTCAGTGGCAGTGGATCAGGAACACAATATTTCTCTCAAGATCAAC
 670 680 690 700 710 720

S L Q P E D F G S Y Y C Q H F W S T P R
 AGCCTGCAACCTGAAGATTTTGGGAGTTATTACTGTCAACATTTTGGAGTACTCCTCGG
 730 740 750 760 770 780

T F G G G T K L E I K R E O K L I S E E
 ACGTTCCGGTGGAGGGACCAAGCTCCGAGATCAAACGGGAACAAAACTCATCTCAGAAGAG
 790 800 810 820 830 840

XhoI

D L N * *
 GATCTGAATTAATAATGATCAAACGGTAATAAGGATCCAGCTCGAATTC
 850 860 870 880

EcoRI

Fig. 6.

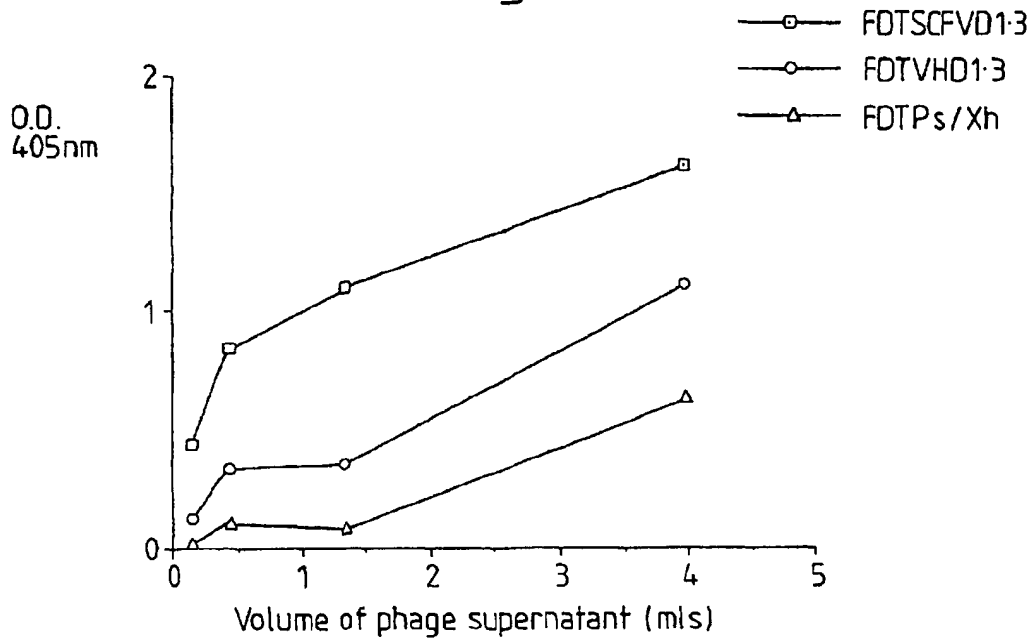


Fig. 7.

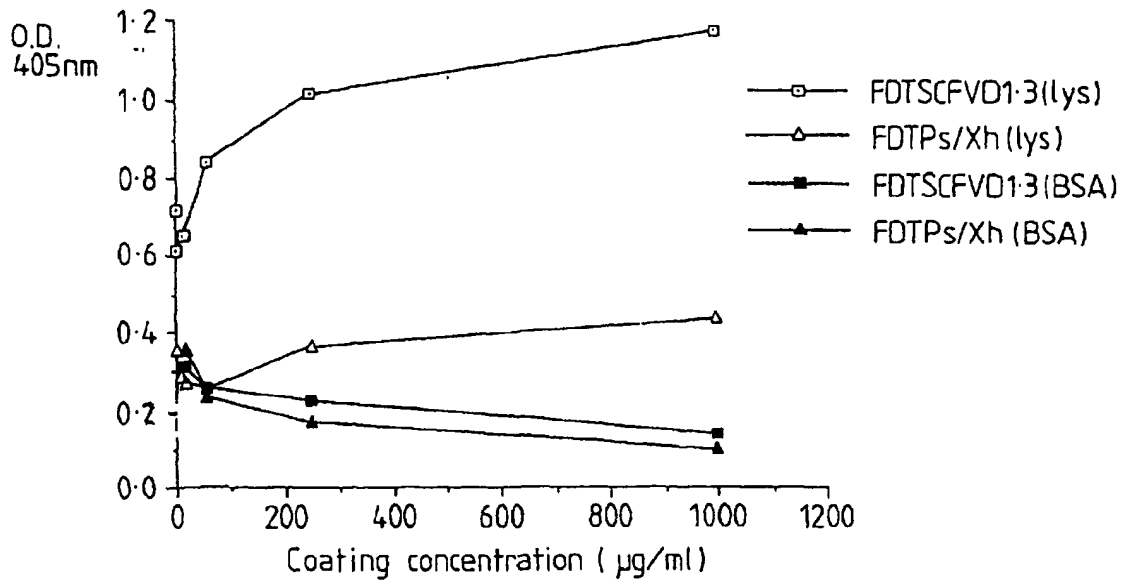


Fig.10.

M K Y L L P T A A
 GCATGCAAATTCTATTTCAAGGAGACAGTCATAATGAAATACCTATTGCCTACGGCAGCC
 10 20 30 40 50 60

A G L L L L A A Q P A M A Q V Q L Q E S
 GCTGGATTGTTATTACTGCTGCCCCAACCAGCGATGGCCCCAGGTGCAGCTGCAGGAGTCA
 70 80 90 100 110 120

G P G L V A P S Q S L S I T C T V S G F
 GGACCTGGCCTGGTGGCGCCCTCAGAGAGCCTGTCCATCACATGCACCGTCTCAGGGTTC
 130 140 150 160 170 180

S L T G Y G V N W V R Q P P G K G L E W
 TCATTAAACCGCTATGGTGTAACCTGGGTTCCGCCAGCCTCCAGGAAGGGTCTCGAGTGG
 190 200 210 220 230 240

L G M I W G D G N T D Y N S A L K S R L
 CTGGGAATGATTTGGGGTGATGCAAACACAGACTATAATTACAGCTCTCAAAATCCAGACTG
 250 260 270 280 290 300

S I S K D N S K S Q V F L K M N S L H T
 AGCATCAGCAAGGACAACCTCCAAGAGCCAAAGTTTCTTAAAAATGAACAGTCTGCACACT
 310 320 330 340 350 360

D D T A R Y Y C A R E R D Y R L D Y W G
 GATGACACAGCCAGGTACTACTGTGCCAGAGAGAGAGATTATAGGCTTGACTACTGGGGC
 370 380 390 400 410 420

Q G T T V T V S S A S T K G P S V F P L
 CAAGGCACCAAGGTACCGTCTCCTCAGCCTCCACCAAGGGCCCATCGGTCTTCCCCCTG
 430 440 450 460 470 480

A P S S K S T S G G T A A L G C L V K D
 GCACCTCTCTCAAGAGCACCTCTGGGGGCACAGCGGCCCTGGGCTGCCTGGTCAAGGAC
 490 500 510 520 530 540

Fig.10 cont.(1)

Y F P E P V T V S W N S G A L T S G V H
TACTTCCCCGAACCGGTGACGGTGTGCTGGAACCTACGGCGCCCTGACCAGCGGCGTGCAC
550 560 570 580 590 600

T F P A V L Q S S G L Y S L S S V V T V
ACCTTCCCGCTGTCTTACAGTCTCTAGGACTCTACTCCCTCAGCAGCGTGGTGACCGTG
610 620 630 640 650 660

P S S S L G T Q T Y I C N V N H K P S N
CCCTCCAGCAGCTTGGGCACCCAGACCTACATCTGCAACGTGAATCACAAGCCCAGCAAC
670 680 690 700 710 720

T K V D K K V E P K S S * *
ACCAAGGTGACAAGAAAGTTCAGCCCCAAATCTTCATAATAACCCGGGAGCTTGCATGCA
730 740 750 760 770 780

M K Y L L P T A A A G L
AATTCTATTTCAGGAGACAGTCATAATGAATACCTATTGCTTACGGCAGCCGCTGGAT
790 800 810 820 830 840

L L L A A Q P A M A D I E L T Q S P A S
TGTTATTACTGCTGCCCCAACCAGCGATGGCCGACATCGAGCTCACCCAGTCTCCAGCCT
850 860 870 880 890 900

L S A S V G E T V T I T C R A S G N I H
CCCTTTCTGGTCTGTGGGAGAACTGTACCATCACATGTGAGCAAGTGGGAATATTC
910 920 930 940 950 960

N Y L A W Y Q Q K Q G K S P Q L L V Y Y
ACAATTATTTAGCATGGTATCAGCAGAAACAGGGAAAATCTCCTCAGCTCCTGGTCTATT
970 980 990 1000 1010 1020

Fig.10 cont.(2)

T T T L A D G V P S R F S G S G S G T Q
 ATACAACAACCTTAGCAGATGGTGTGCCATCAAGGTTCAAGTGGCAGTGGATCAGGAACAC
 1030 1040 1050 1060 1070 1080

Y S L K I N S L Q P E D F G S Y Y C Q H
 AATATTCTCTCAAGATCAACAGCCTGCAGCCTGAAGATTTTGGGAGTTATTACTGTCAAC
 1090 1100 1110 1120 1130 1140

F W S T P R T F G G G T K L E I K R T V
 ATTTTTGGAGTACTCCTCGGACGTTGGTGGAGGCACCAAGCTCGAGATCAAACCGACTG
 1150 1160 1170 1180 1190 1200

A A P S V F I F P P S D E Q L K S G T A
 TGGCTGCACCATCTGTCTTCATCTTCCCGCCATCTGATGAGCAGTTGAAATCTGGAACG
 1210 1220 1230 1240 1250 1260

S V V C L L N N F Y P R E A K V Q W K V
 CCTCTGTTGTGTGCGCTGCTGAATAACTTCTATCCCAGAGAGGCCAAAGTACAGTGGAAAG
 1270 1280 1290 1300 1310 1320

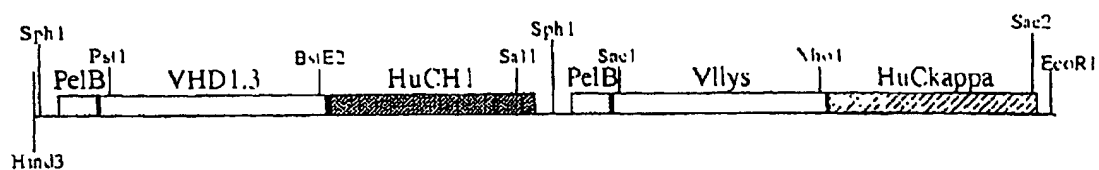
D N A L Q S G N S Q E S V T E Q D S K D
 TGGATAACGCCCTCCCAATCGGGTAACTCCAGGAGAGTGTACAGAGCAGGACAGCAAGG
 1330 1340 1350 1360 1370 1380

S T Y S L S S T L T L S K A D Y E K H K
 ACAGCACCTACAGCCTCAGCAGCACCTGACGCTGAGCAAGCAGACTACGAGAAACACA
 1390 1400 1410 1420 1430 1440

V Y A C E V T H Q G L S S P V T K S F N
 AAGTCTACGCCCTGGGAGTCAACCATCAGGGCCTGAGCTCGCCCGTCACAAAGAGCTTCA
 1450 1460 1470 1480 1490 1500

R G E S * *
 ACCGCGGAGAGTCATAGTAAGAATTC
 1510 1520

Fig. 10 cont. (3)



FabD1.3 in pUC19

Fig. 11.

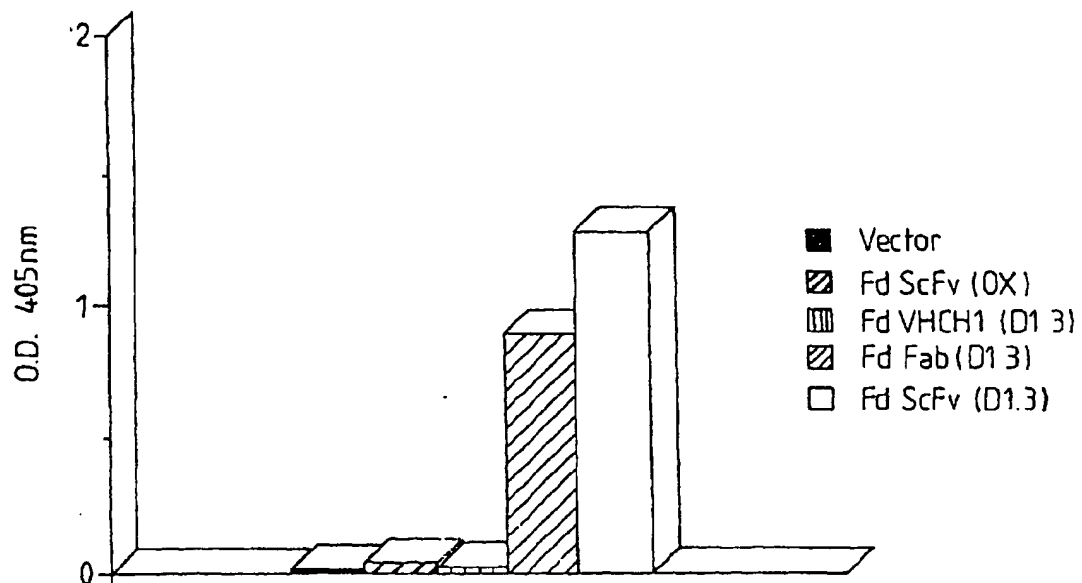


Fig.12.

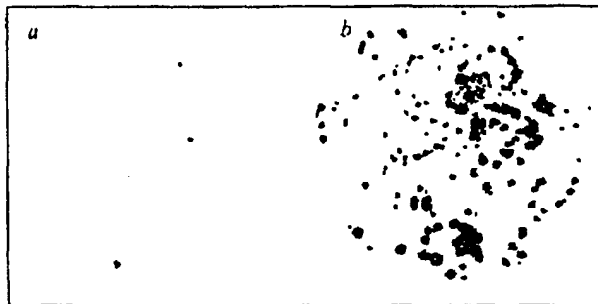


Fig. 13.

Q V Q L Q E S G G L V Q P G G
CAG GTG CAG CTG CAG GAG TCA GGA GGA GGC TTG GTA CAG CCT GGG GGT
PstI
S L R L S C A T S G F T F S N Y
TCT CTG AGA CTC TCC TGT GCA ACT TCT GGG TTC ACC TTC AGT AAT TAC
Y M G W V R Q P P G K A L E W L
TAC ATG GGC TGG GTC CGC CAG CCT CCA GGA AAG GCA CTT GAG TGG TTG
G S V R N K V N G Y T T E Y S A
GGT TCT GTT AGA AAC AAA GTT AAT GGT TAC ACA ACA GAG TAC AGT GCA
S V K G R F T I S R D N F Q S I
TCT GTG AAG GGG CGG TTC ACC ATC TCC AGA GAT AAT TTC CAA AGC ATC
L Y L Q I N T L R T E D S A T Y
CTC TAT CTT CAA ATA AAC ACC CTG AGA ACT GAG GAC AGT GCC ACT TAT
Y C A R G Y D Y G A W F A Y W G
TAC TGT GCA AGA GGC TAT GAT TAC GGG GCC TGG TTT GCT TAC TGG GGC
Q G T L V T v s s g g g g s g g g g s
CAA GGG ACC CTG GTC ACC gtc tcc tca ggtaggagcggttcaggcggagggtggetct
BstEII
g g g g s d i E L T Q T P L S L P V
ggcggtggcggtcgac atc GAG CTC ACC CAA ACT CCA CTC TCC CTG CCT GTC
SacI
S L G D Q A S I S C R S S Q S I
AGT CTT GGA GAT CAA GCC TCC ATC TCT TGC AGA TCT AGT CAG AGC ATT
V H S N G N T Y L E W Y L Q K P
GTA CAT AGT AAT GGA AAC ACC TAT TTA GAA TGG TAC CTG CAG AAA CCA
PstI
G Q S P K L L I Y K V S N R F S
GGC CAG TCT CCA AAG CTC CTG ATC TAC AAA GTT TCC AAC CGA TTT TCT
G V P D R F S G S G S G T D F T
GGG GTC CCA GAC AGG TTC AGT GGC AGT GGA TCG GGG ACA GAT TTC ACA
L K I S R V E A E D L G V Y Y C
CTC AAG ATC AGC AGA GTG GAG GCT GAG GAT CTG GGA GTT TAT TAC TGC
F Q G S H V P Y T F G G G T K L
TTT CAA GGT TCA CAT GTT CCG TAC ACG TTC GGA GGG GGG ACC AAG CTC
E I K R
GAG ATC AAA CGG
XhoI

Fig. 14.

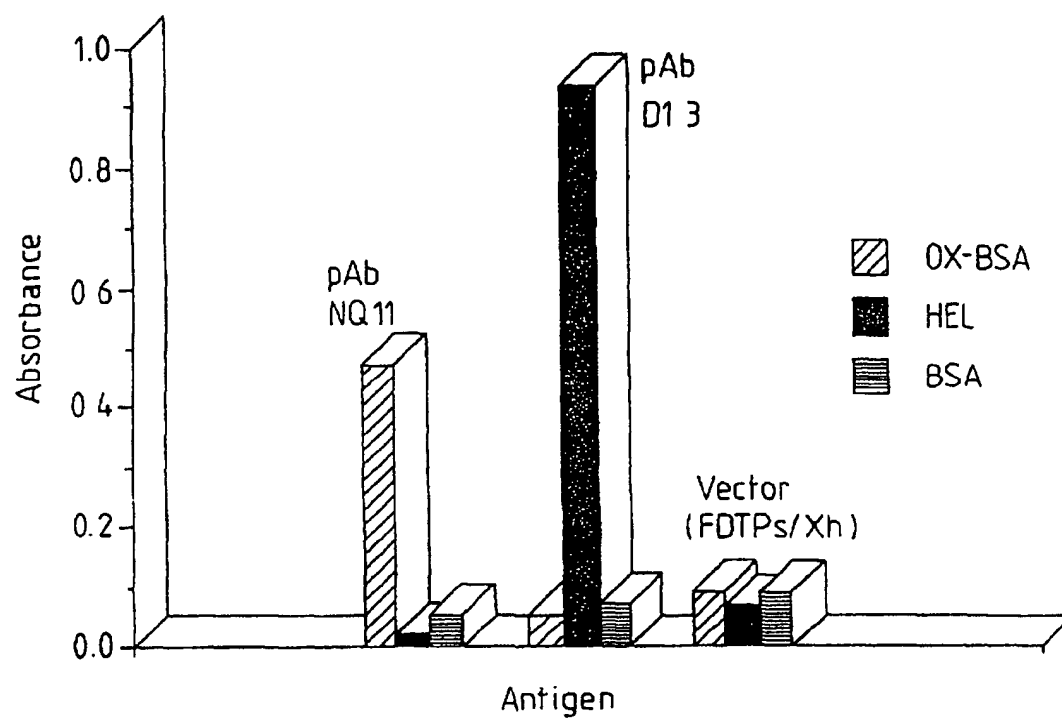


Fig. 15.

5' END

TCT CAC AGT GCA CAA ACT GTT GAA CGG ACA CCA GAA ATG CCT GTT CTG
 ApaI

3' END

K A A L G L K
 AAA GCC GCT CTG GGG CTG AAA GCG GCC GCA GAA ACT GTT GAA AGT etc
 Not I

Fig.16(1)

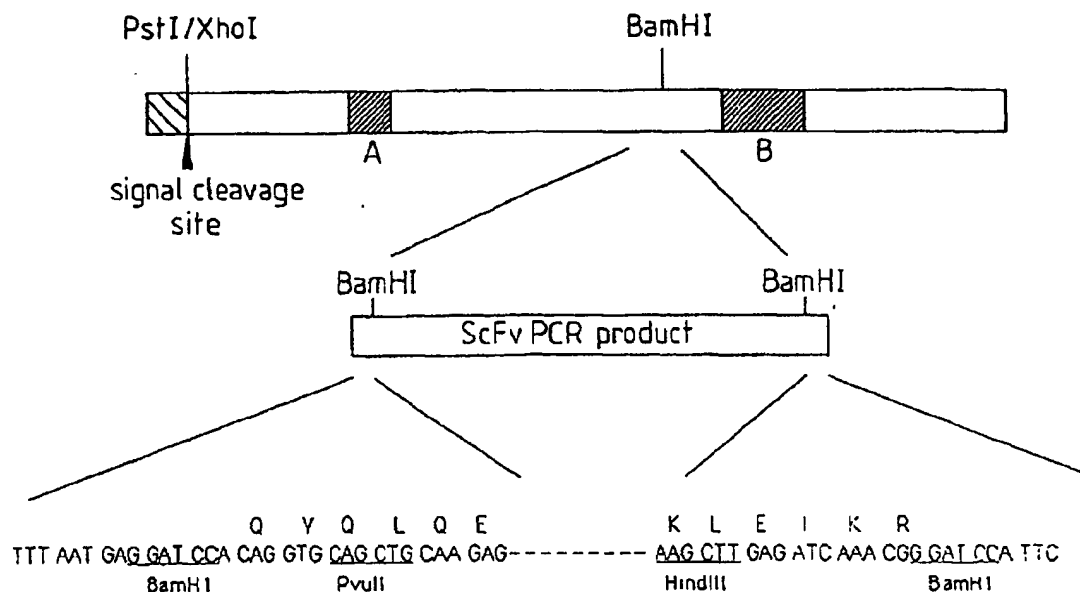


Fig.16(2)

A (1834) 5' GAG GGT GGT GGC TCT
 " " "C " "
 " " "C " "
 " " "C " ACT 3'(1839)

B (2284) 5' - GGC GGC GGC TCT
 - GGT GGT GGT "
 - " GGC GGC "
 GAG " " GGC "
 " " " GGT "
 " " " GGC "
 " " " GGT "
 - " " GGC " 3'(2379)

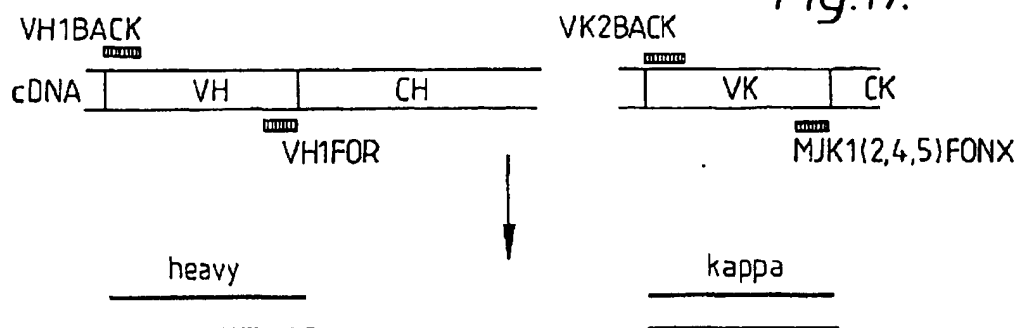
Reverse complement of mutagenic
 oligo 63Bamlink

5' GAG GGT GGC GGA TCC

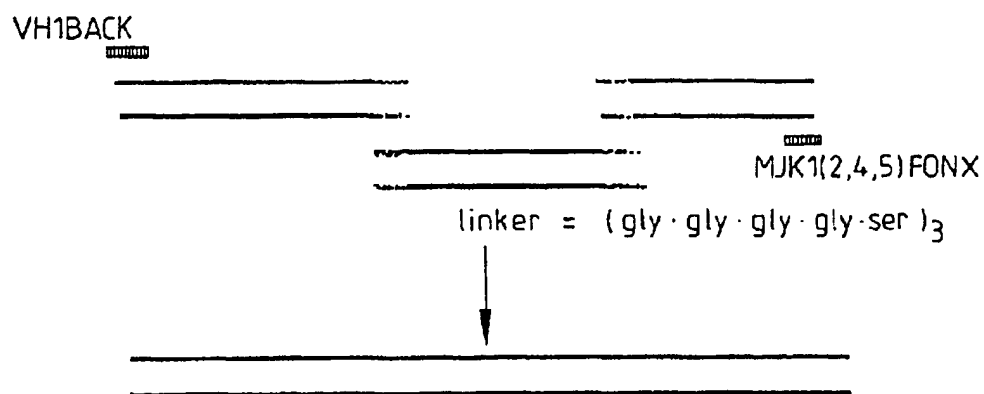
GAG GGT GGC GG 3'

Fig.17.

1) PRIMARY PCR



2) ASSEMBLY PCR



3) ADDING RESTRICTION SITES

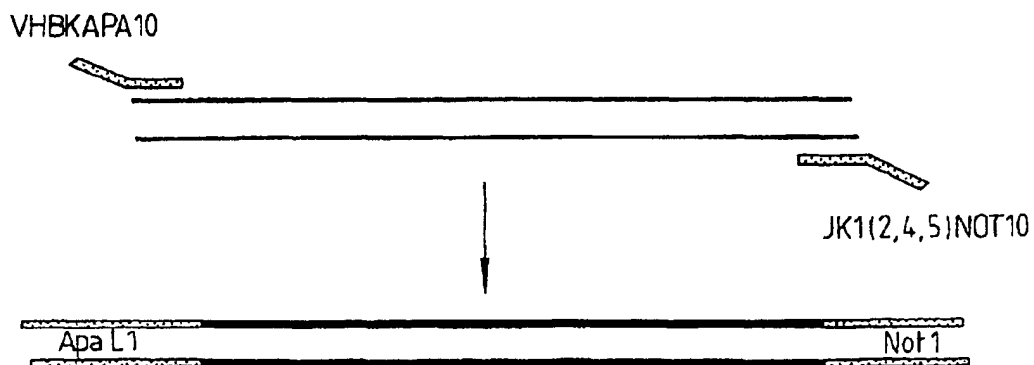


Fig.18.

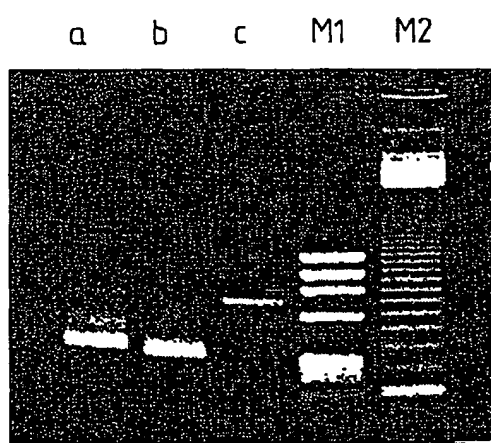


Fig. 19

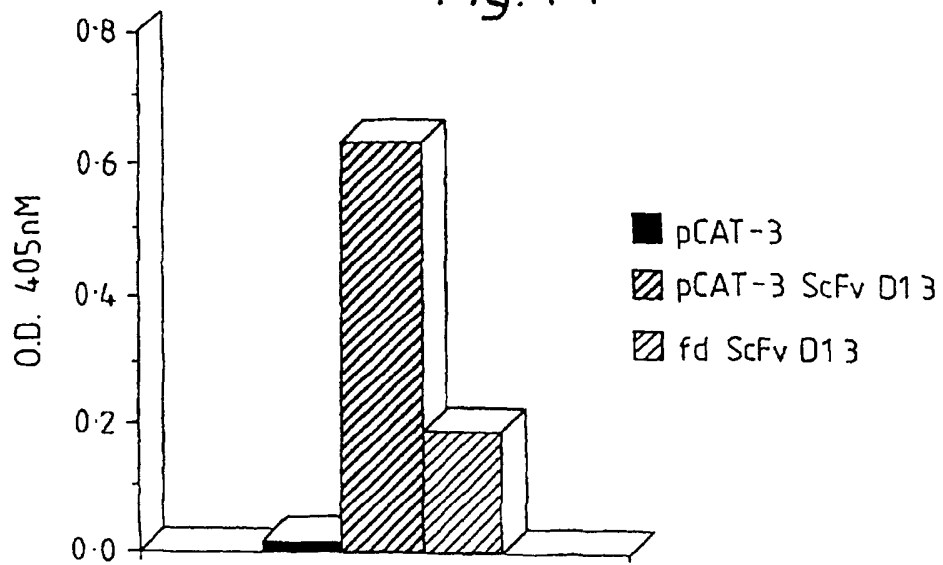
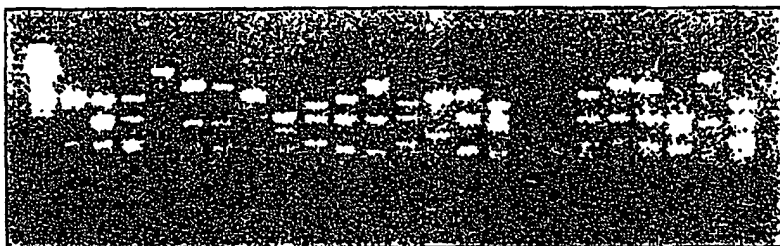


Fig. 20

d

M



M

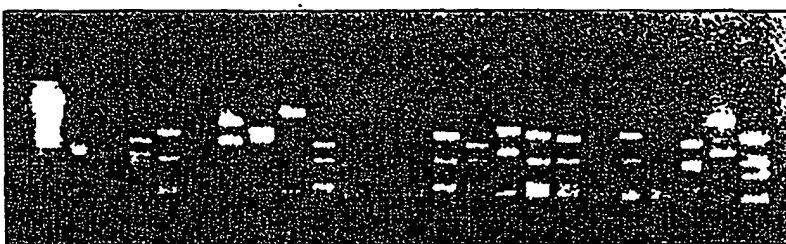


Fig. 21

VH sequences

from combinatorial library:

	CDR1	CDR2	CDR3	
A	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x4 1
B	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x9 1
C	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x3 1
D	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x3 1
E	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x3 1
F	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x3 1
G	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x3 1
H	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x3 1

from hierarchical library VH-rep x V_k-d:

I	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1
J	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1
K	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x3 1
L	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x3 1
M	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1
N	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1
O	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1
P	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1
Q	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1
R	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x3 1
S	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 x6 1
T	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1
U	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1
V	QVQLQSGQELARPQASVKNSSCKASGYTFT	WYQRPQQGLEWIG	KATLTADKSSSTAYHQLSBLTSEDNAVYYCAR	WQQTITVTVS8 1

Fig. 2/cont.

V κ sequences

from combinatorial library:

	CDR1	CDR2	CDR3		
a	DIELTQSPALNSASPOEKVTMT	LAAGTLES	LOVASTPT	FOAGTKLEIKRA	V
b	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	IV
c	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	IV
d	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	IV
e	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	IV
f	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	IV
g	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	IV

from hierarchical library VH-B x V κ -rep:

h	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	IV/VI
i	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
j	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
k	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
l	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
m	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
n	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
o	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
p	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
q	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
r	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
s	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
t	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
u	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
v	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V
w	DIELTQSPALNSASPOEKVTMT	STSNLAS	QOYSDYPLT	FOAGTKLEIKRA	V

Fig. 22

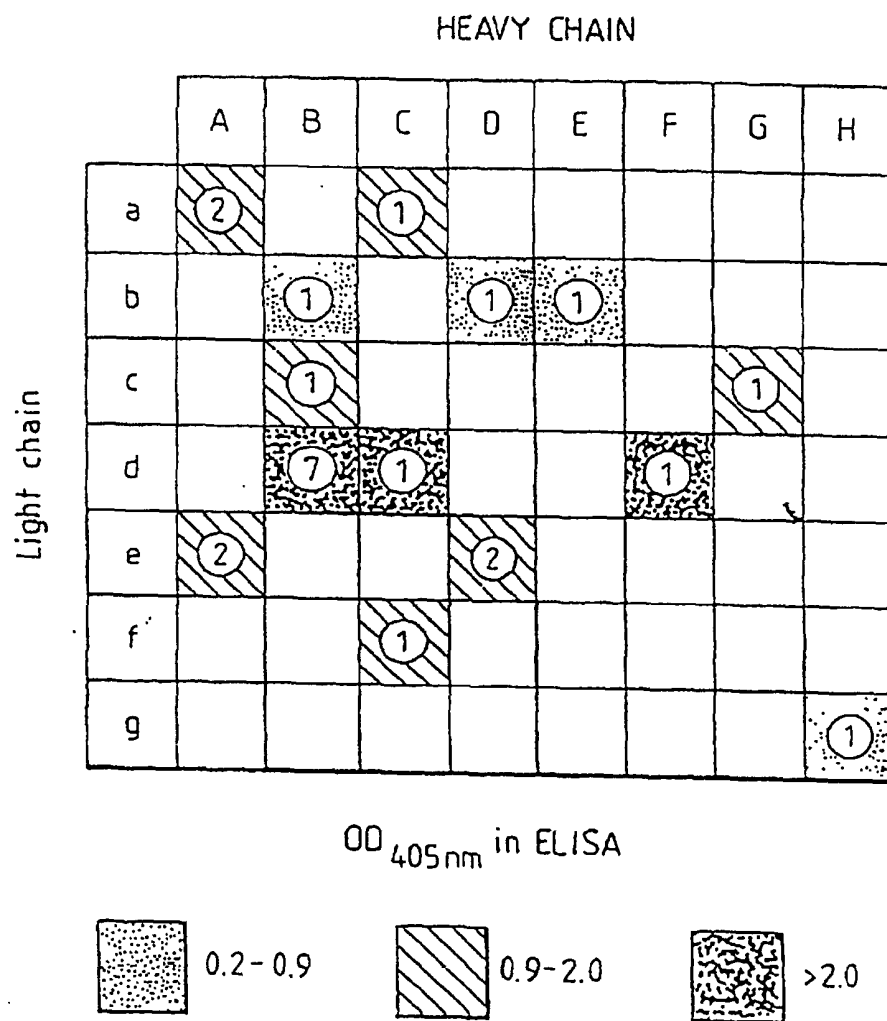


Fig. 23(a)

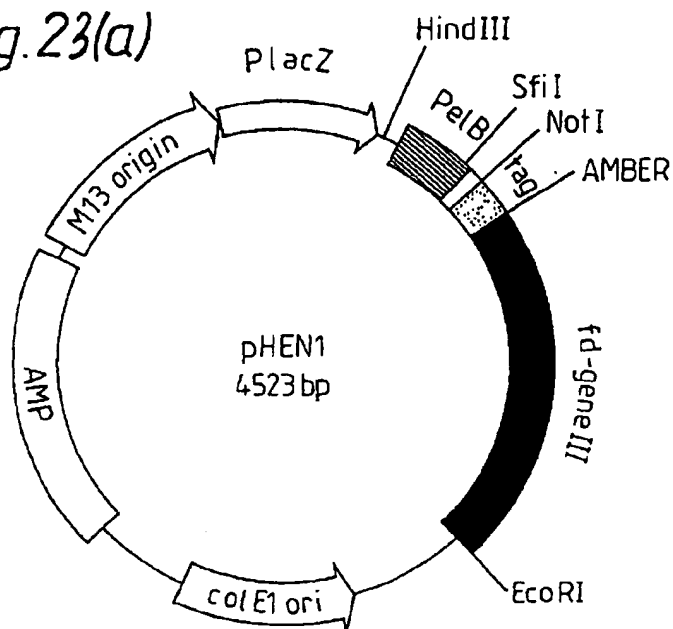


Fig. 23(b)

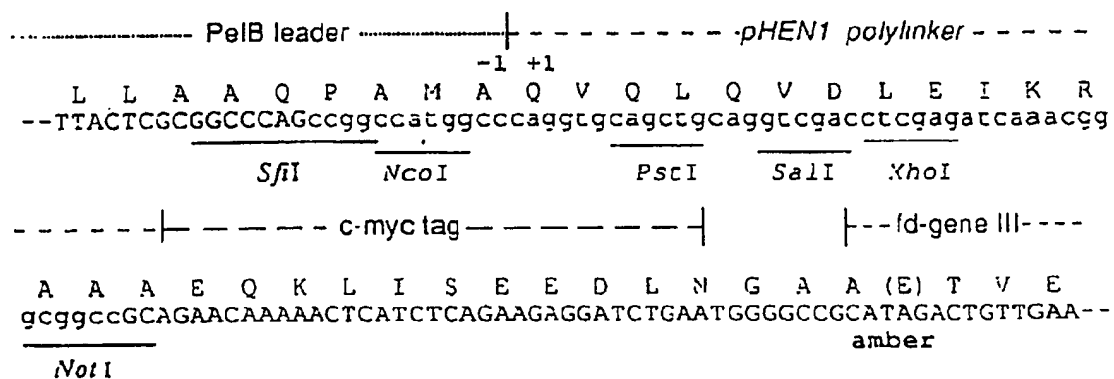


Fig. 24

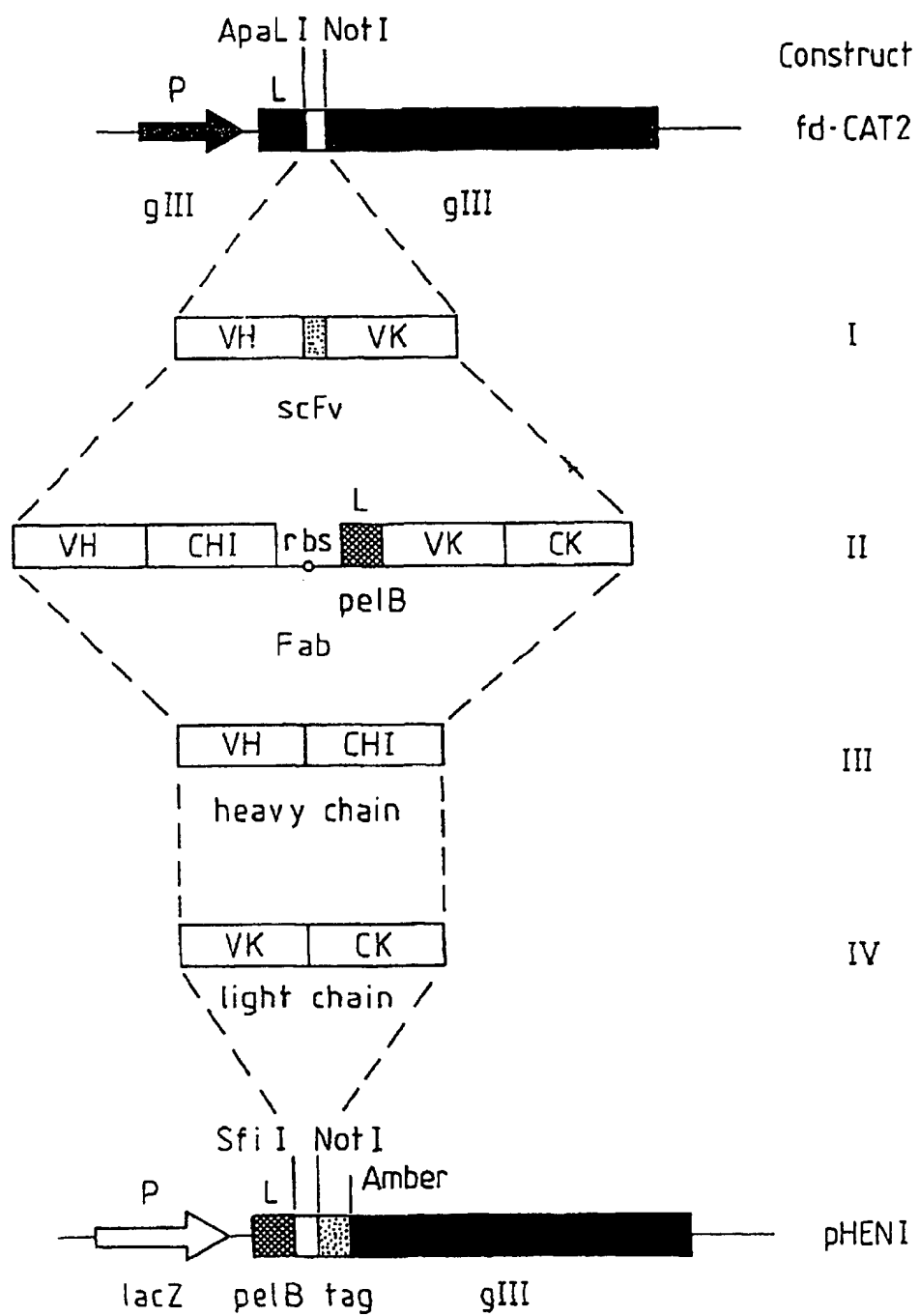


Fig. 25

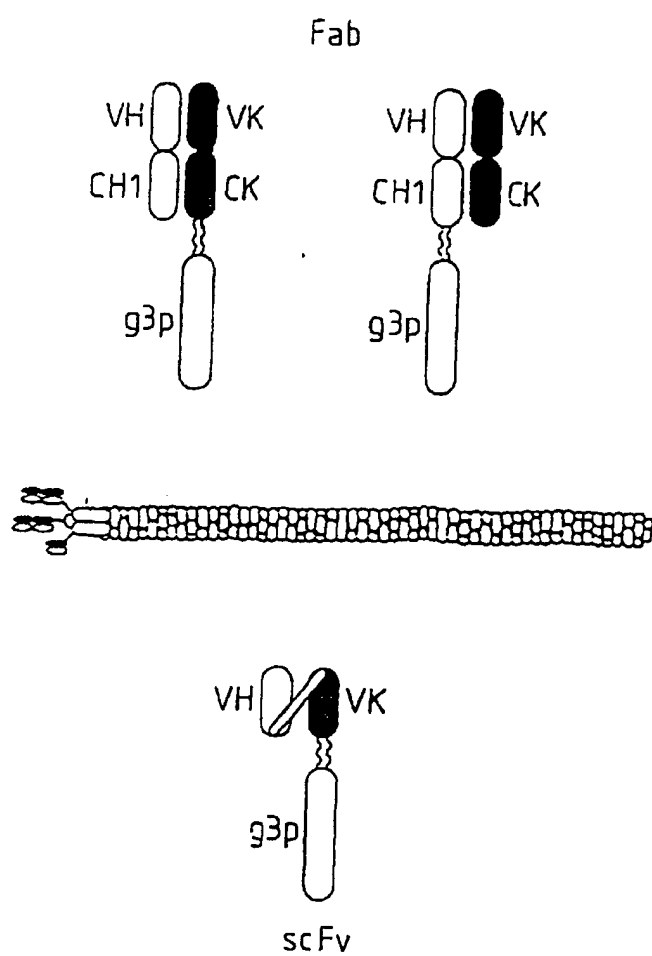


Fig. 26

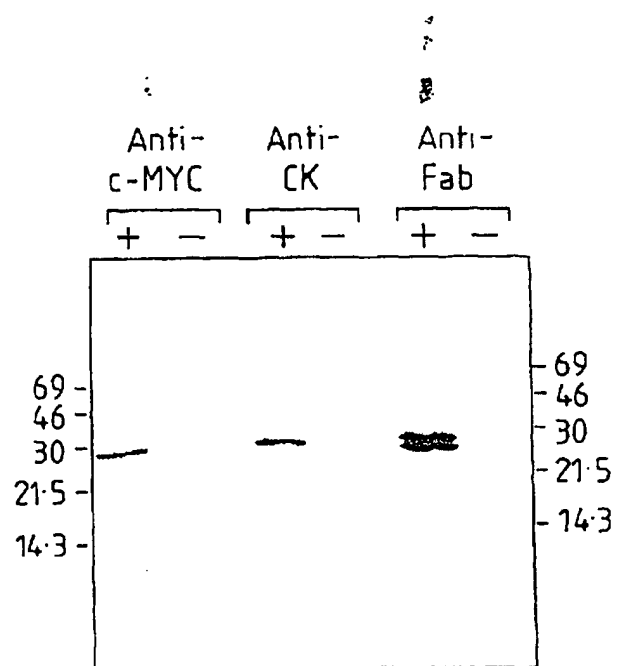


Fig. 27

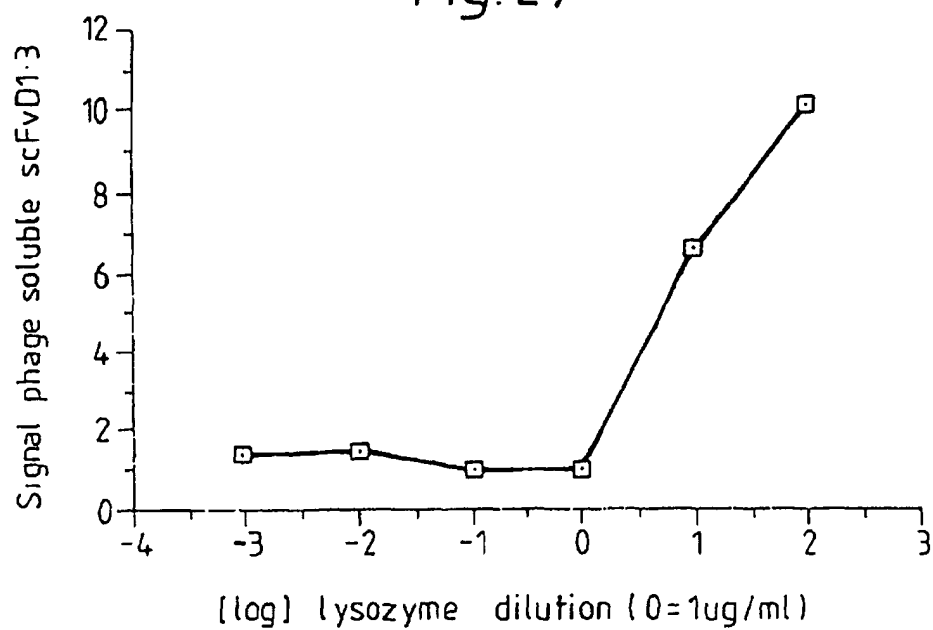


Fig. 28

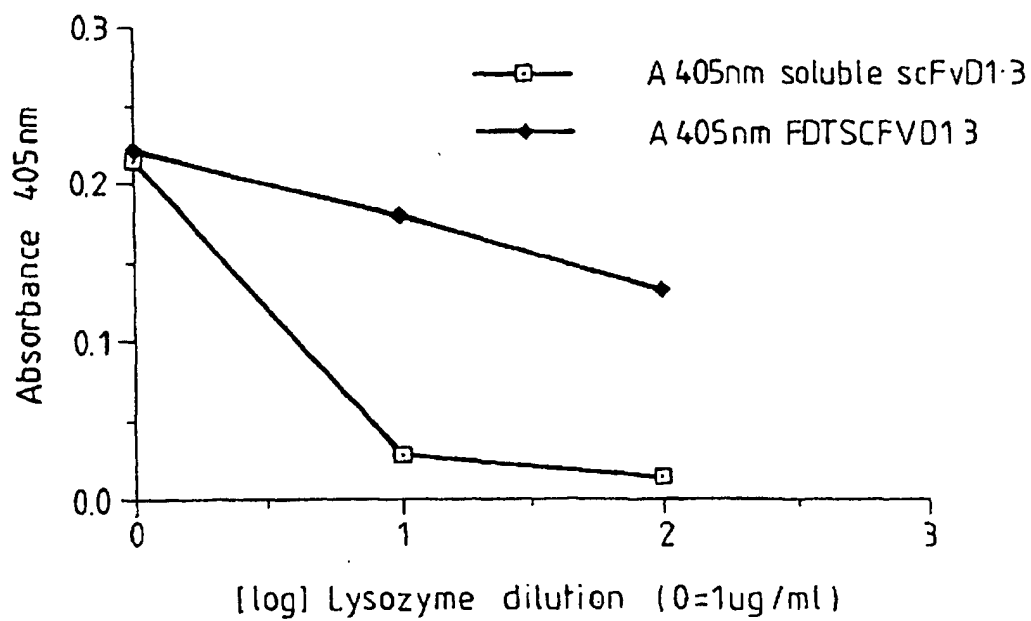


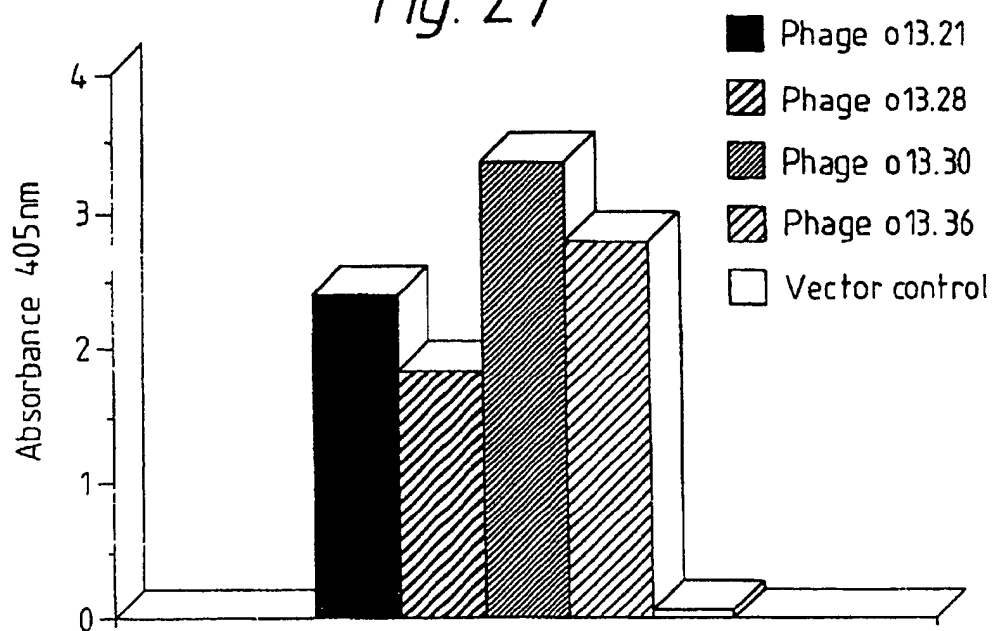
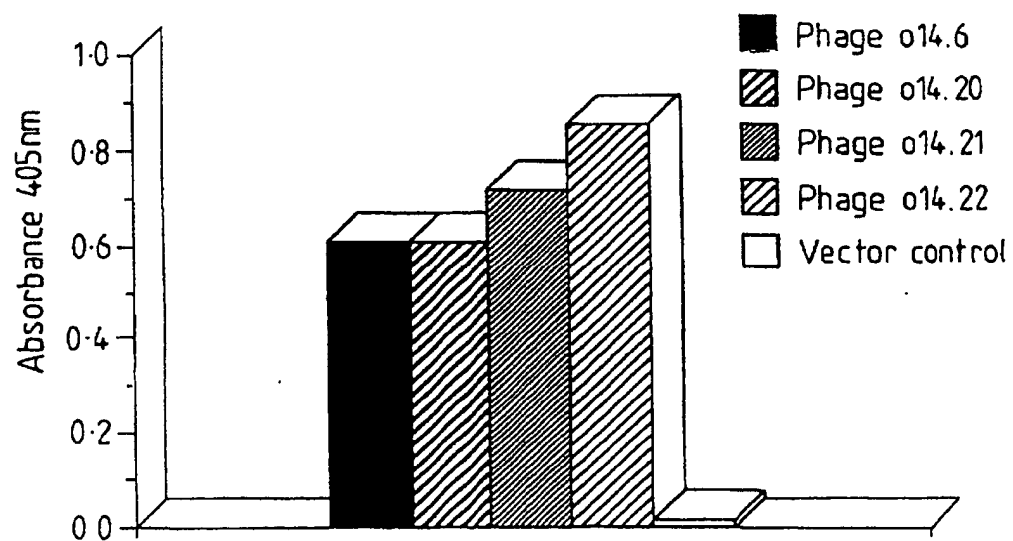
Fig. 29*Fig. 30*

Fig. 31

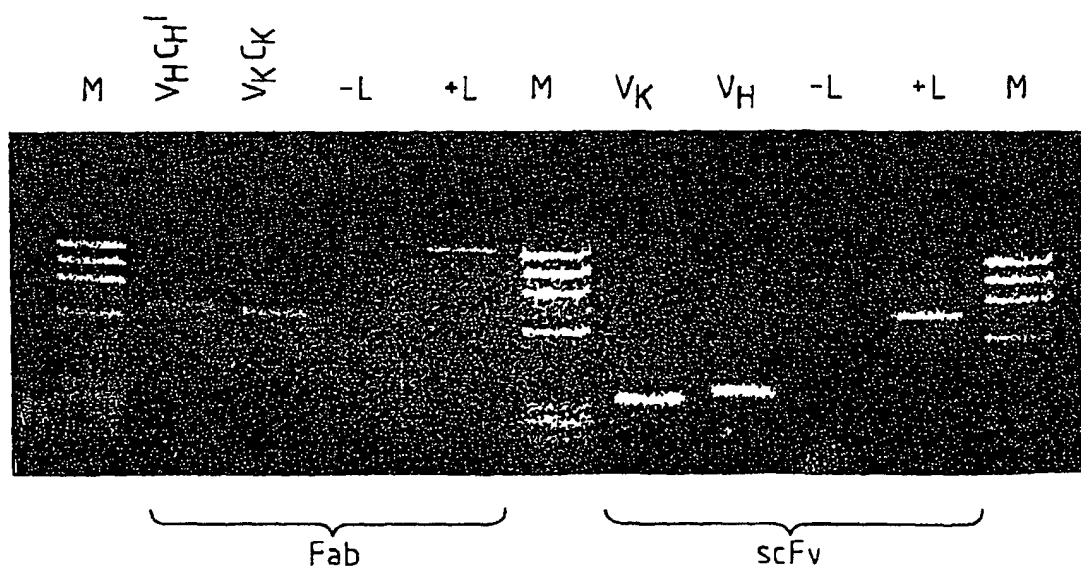


Fig. 32

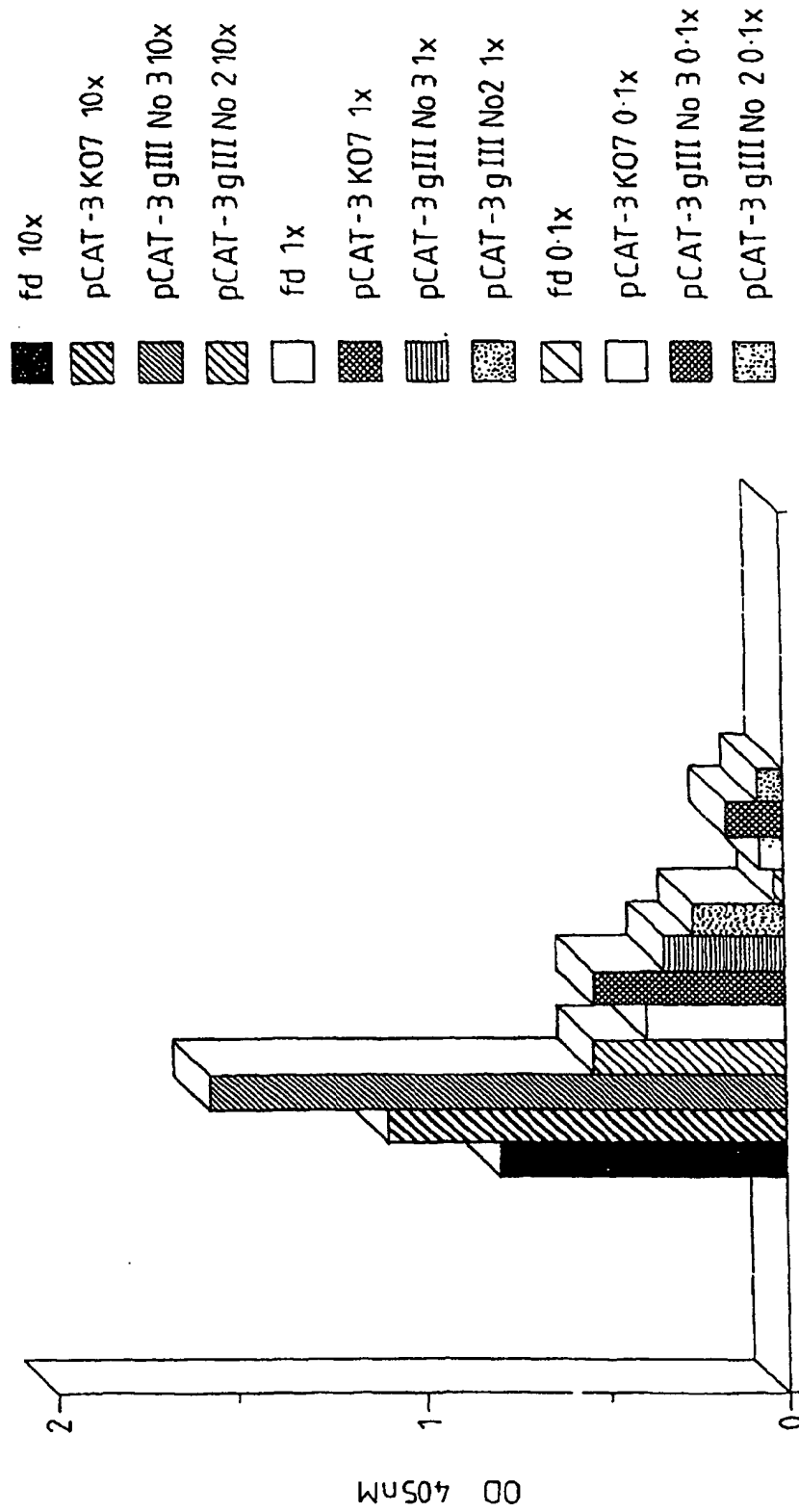


Fig.33

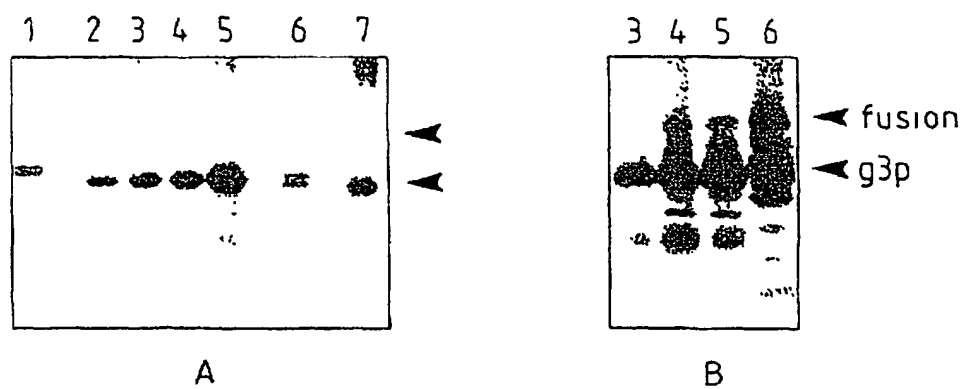


Fig. 34

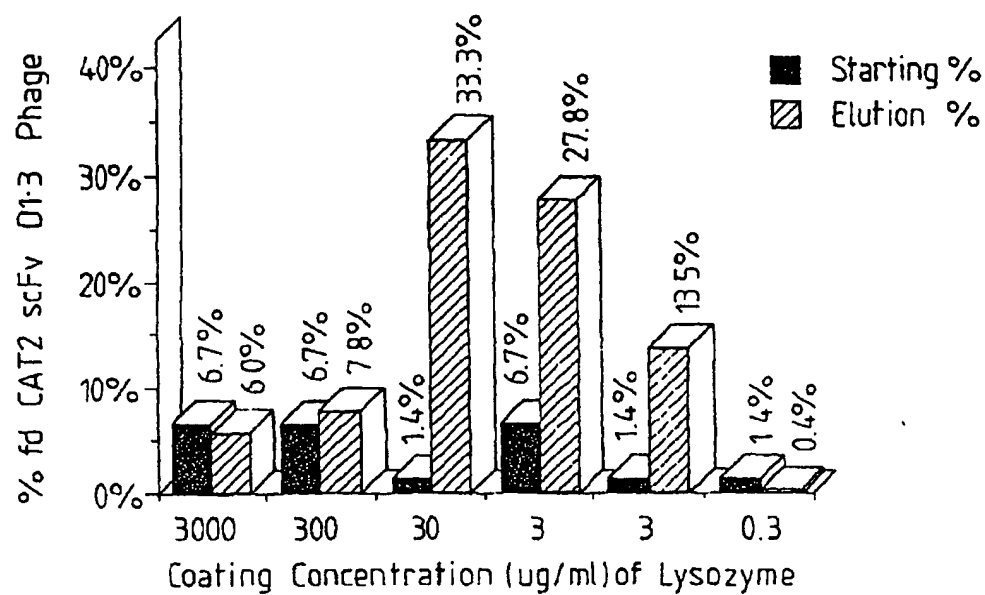


Fig. 35

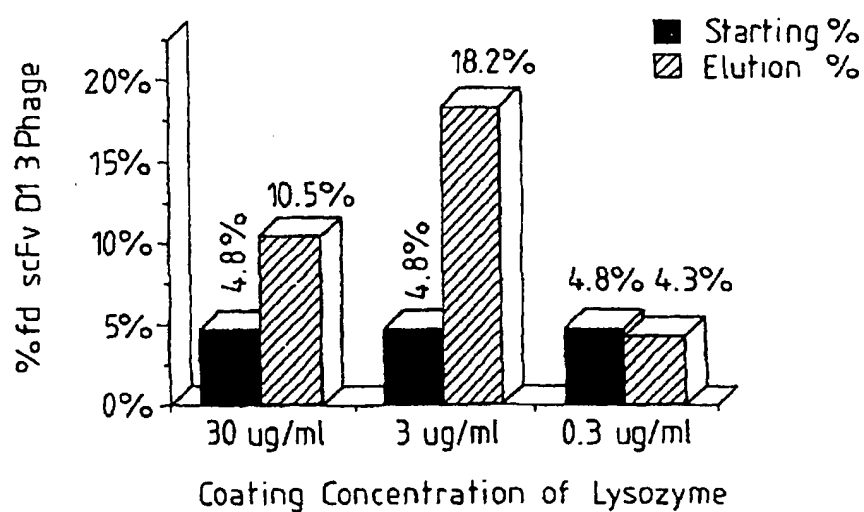


Fig. 36 (i)

10	20	30	40	50	60	70	80	90
TTCTATTCTACAGTGCACAGGTC	CGAGTCCAGGTCGACGTCGTGGGCT	GTGACCTTGTCAAGCCTGGGGCT	TCAGTGAAGCTGTCTCTGCAAGGCT					
AAGATAAGAGTGTACAGGTGTCCAGGT	CGACGTCGTGACACCCGAGCTCGAACAC	TCTCGGACCCCGAAGTCACTTCGACAC	CGACGTCGACACGACGTCCTCCGA					
PhETyrSerHisSerAlaGlnValGlnLeu	GlnSerGlyAlaGluLeuVallysProGly	AlaSerValLysLeuSerCysLysAla						
100	110	120	130	140	150	160	170	180
TTCTGGCTACACCTTACACAGCTACT	GGATGCACCTGGGTGAAGCAGAGGCT	TGGACGAGCCCTTGGAGTGGATTTGG	AMGGATTGNTCTCTAAT					
AGACCGATGTGGAAAGTGGTCGATGAC	CTTACCTTCGTCCTCGGACCTTGC	TCCGGACCTTGC	TCCGGAACTCACCTTAACCTTCTCA					
AGATTAAGTGGTGGTGGTGGTGGTGGT	GGTGGTGGTGGTGGTGGTGGTGGT	GGTGGTGGTGGTGGTGGTGGTGGT	GGTGGTGGTGGTGGTGGTGGTGGT					
SerGlyTyrThrPheThrSerTyrTrpMet	HisTyrValLysGlnArgProGlyArgGly	LeuGluTrpLleGlyArgLleAspProAsn						
190	200	210	220	230	240	250	260	270
AGTGGTGGTACTAAGTACAATGAGAGT	TCAAGGCAAGGCCACACCTGAC	TGTAGACAAACCTCCAGCACACGCT	ACATACATGCAGCTCAGC					
TCACCAACCATGATTGATGATGATGAT	GATGATGATGATGATGATGATGAT	GATGATGATGATGATGATGATGAT	GATGATGATGATGATGATGATGAT					
SerGlyGlyThrLysTyrAsnGluLysPhe	LysSerLysAlaThrLeuThrValAspLys	ProSerSerThrAlaTyrMetGlnLeuSer						
280	290	300	310	320	330	340	350	360
AGCCTGACATCTGAGGACTCTGGGCT	TATTATTGTGGAAGTACGACTACGGT	TAGTACGCTACTTTGACTACTTGGG	CCCAAGGGAC					
TCGGACTGTAGACTCTGAGACGCCAG	ATAATAACACGTTCTATGCTGATGCC	ATCATCGATGATGAAC	TGATGACCCCGGTTCCCTGG					
SerLeuThrSerGluAspSerAlaValTyr	TyrCysAlaArgTyrAspTyrGlySer	SerTyrPheAspTyrTrpGlyGlnGlyThr						
370	380	390	400	410	420	430	440	450
ACGGTCACCGTCTCCTCAGGTGGAGG	CGGTTACGGCGGAGGTGGCTCTGG	CGGTGGCGGATCCAGGCTGTTGGG	ACACAGGAATCTGCA					
TGCCAGTGGCAGAGGATCCACCTCCG	CCCAAGTCCCGCTCCACCGAGACCG	CCACCGCTAGGTCGACACACCTGT	GTCCTTAGACGT					
ThrValThrValSerSerGlyGlyGlySer	GlyGlyGlySerGlyGlyGlySerGly	GlyGlyGlySerGlnAlaValGlyThr	GlnGluSerAla					
460	470	480	490	500	510	520	530	540
CTCACCAATCACCCTGGTGAAACAGT	CACACTCTGTCGCTCAGTACTGGG	CGTGTACAACTATGCCAACTGGGT	TCCAA					
GAGTGGTGTAGTGACCACTTTGTCA	GTTGAGTGAAACACCGAGTTCATG	ACCCCGACAAATTTGATCATTGAT	CACGGTTGACCCAGGTT					
LeuThrThrSerProGlyGluThrValThr	LeuThrCysArgSerSerThrGlyAla	ValThrSerAsnTyrAlaAsnTrpValGln						
550	560	570	580	590	600	610	620	630
GAATAACAGATCATTTATTCACTGGT	CTAATAGGTGTACCAACACCGAGCT	TCCAGGTGTTCTGCCAGATTCTCA	GGCTCCCTTGATT					
CTTTTGGTCTAGTAAATAAGTGACCA	GATTATTCACCATGTTGTTGGTCC	GAGTCCACAAAGGACGGTCTAAG	AGTCCGAGGAGCTAA					
GluLysProAspHisLeuPheThrGlyLeu	lleGlyGlyThrAsnAsnArgAlaProGly	ValProAlaArgPheSerGlySerLeu	Ile					

Fig. 36(ii)

```

640      650      660      670      680      690      700      710      720
GGAGACAAGGCTGCCCTCACCATCACAGGGGCACAGACTGAGGATGAGGCANTATATTCTGTCTCTATGGTACAGCAACCATTTGGGTG
CCTCTGTTCCGACGGAGTGTAGTGTCCCGTGTCTGACTCCTACTCCGTTATATAAAGACACGAGATACCATGTCTCGTTGGTAACCCAC
GlyAspLysAlaAlaLeuThrIleThrGlyAlaGlnThrGluaspGluAlaIleTyrPheCysAlaLeuTrpTyrnberAsnHisTrpVal

730      740      750      760      770
TTCGGTGGAGGAACCAAACCTGACTGTCTCTCGAGATCAAAACGGGCGGCCGC
AAGCCACCTCCTTGGTTGACTGACAGGAGCTCTAGTTTCCCCGCCGCGC
PheGlvGlyGlyThrLysLeuThrValLeuGluIleLysArgAlaAla

```

Fig. 37

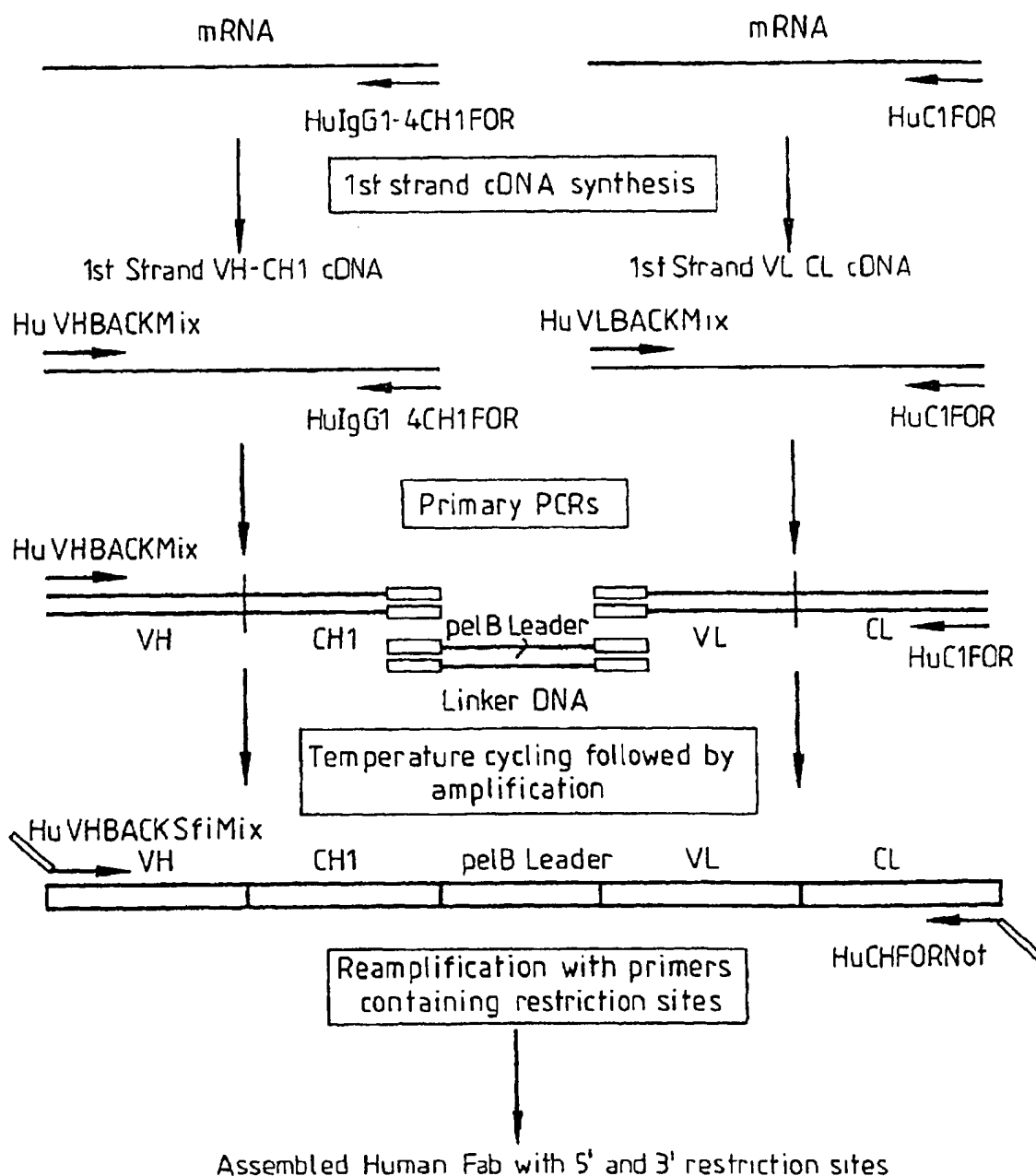
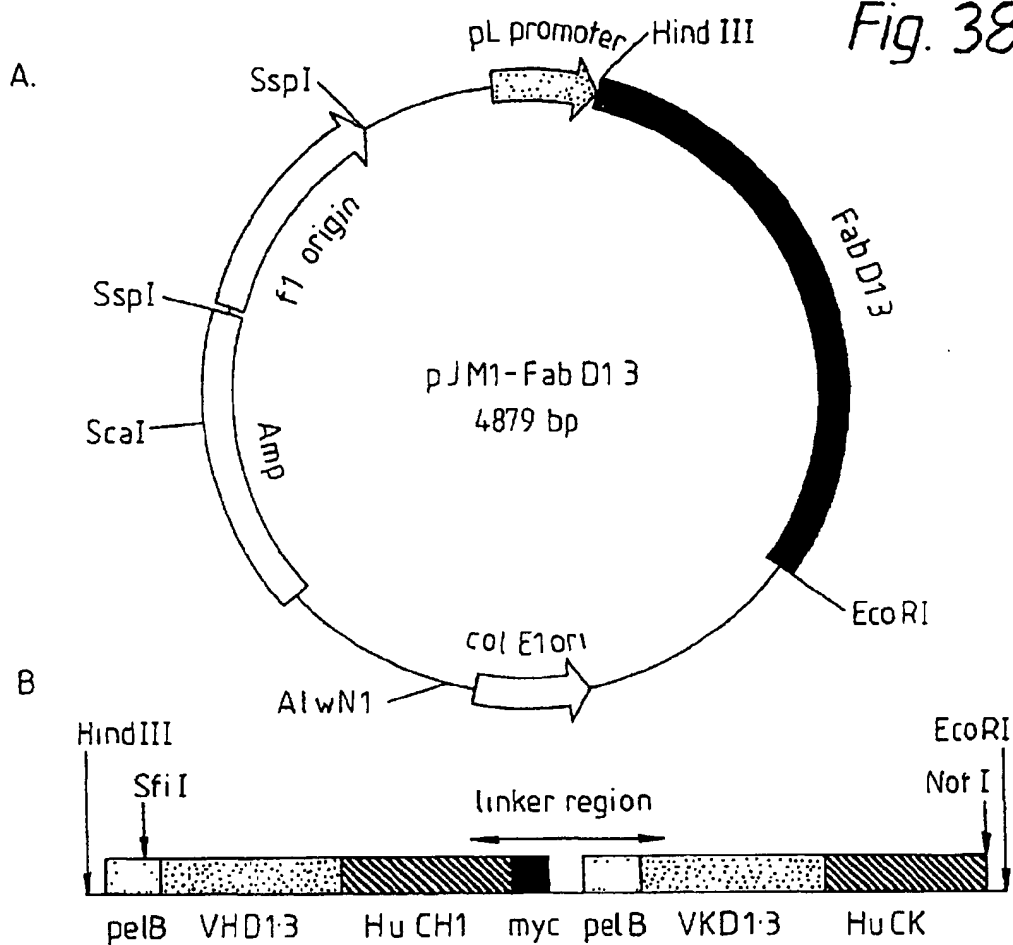


Fig. 38



C. Sequence of linker region

← 3' Human CH1 and hinge →
 K P S N T K V D K K V E P K S S T K T H T
 AACCCGAGCAACACCAAGGTCGACAAGAAAGTTGAGCCCAATCTTCAACTAAGACGCACACA

→ myc peptide tag →
 S G G E Q K L I S E E D L N * *
 TCAGGAGGTGAACAGAAGCTCATCTCAGAAGAGGATCTGAATTAATAAGGGAGCTTGCATGCA

← pelB leader →
 M K Y L L P T A A A G L
 AATTCTATTTCAGGAGACAGTCATAATGAAATACCTATTGCCTACGGCAGCCGCTGGATTGT

→ 5' Vk →
 L L P A A Q P A M A D I E L T Q S P
 TATTACCTGCTGCCCAACCAGCGATGGCGACATCGAGTTCAACCCAGTCTCC

Fig. 39

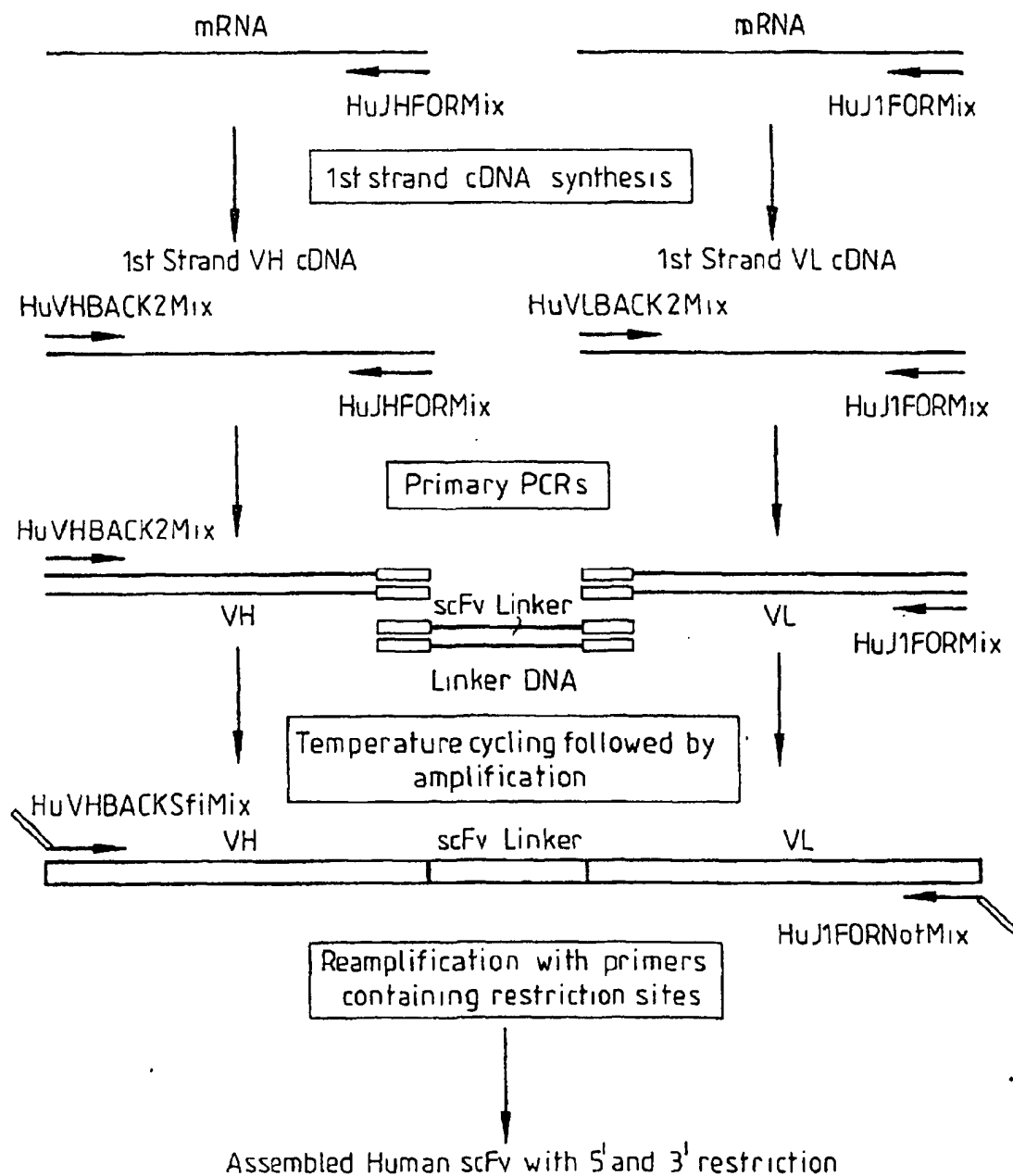


Fig. 40

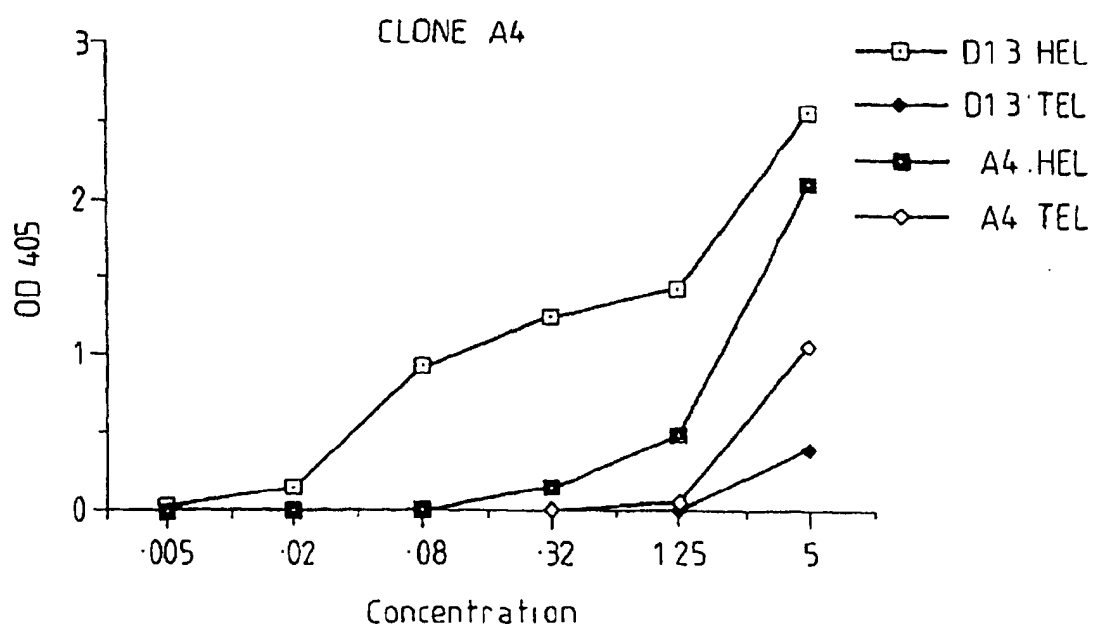
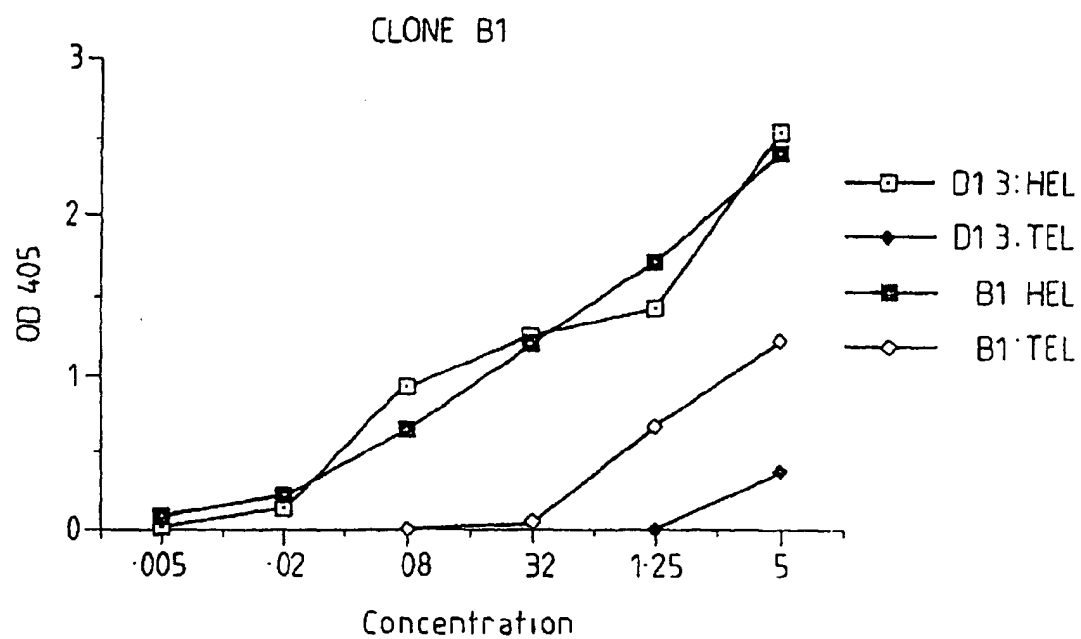


Fig. 41

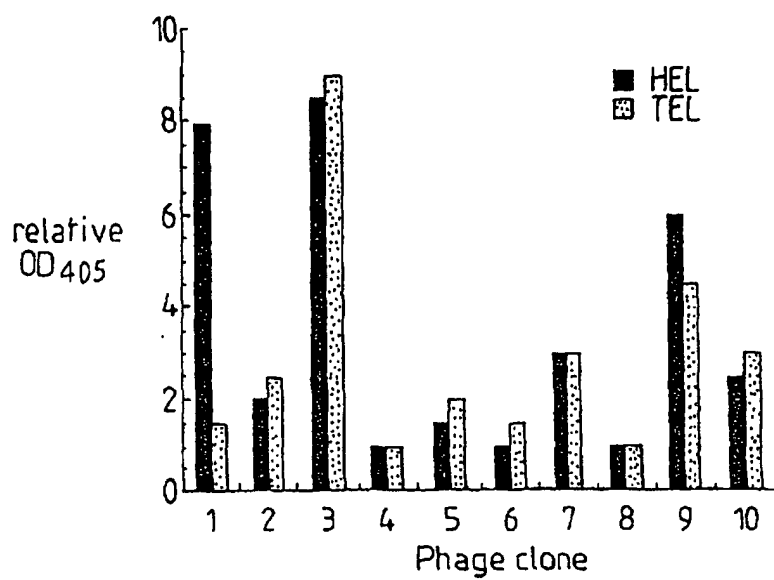


Fig. 43

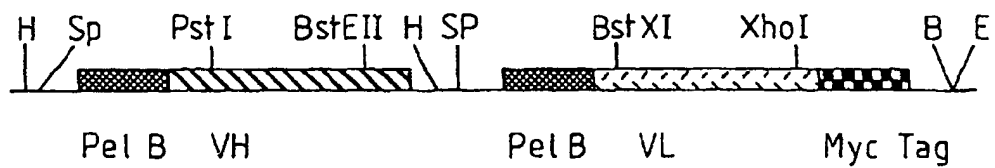


Fig. 42

	CDR 1	CDR 2
D1.3	DIQMTQSPASLSASVGETVTITCRASGNIHNYLA WYQQKQKSPQLLVYYTTLAD	
M1F	DIELTQSPSSLSASLGERVSLTCRASQDIGSSLN WLQQEPDGTIKRLIYATSSLDS	
M21	DIELTQSPALMAASPGEKVTITCSVS88I888NLHWYQQKSETSPKPWIYGT8NL88	
	CDR 3	
D1.3	GVPSRFSGSGGTQYSLKINSLQPEDFGSYQCQHFWSPTPRTFGGTKLEIKR	
M1F	GVPKRFSGRSGSDYSLTISSESEDFVDYVYCLQYASSPWTFFGGTKLELKR	
M21	GVPVRFSGSGGTSYSLTISSEAEADAATYCCQQWSSYPPLTFGAGTKLEIKR	